

FORMPTO-1390  
(REV 12-29-99)

U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

**TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371**

IR 3492NP-PCT

U S APPLICATION NO. (If known, see 37 CFR 1.5)

**09/869251**

INTERNATIONAL APPLICATION NO.

PCT/US00/30953

INTERNATIONAL FILING DATE

09 November 2000 (9.11.00)

PRIORITY DATE CLAIMED

09 November 1999 (9.11.99)

**TITLE OF INVENTION TACK FREE SURFACE CURES OF POLYMERS BY ORGANIC PEROXIDES IN THE PRESENCE OF AIR**

**APPLICANT(S) FOR DO/EO/US ATOFINA Chemicals, Inc., et al.**

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☐ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
  - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☐ has been transmitted by the International Bureau.
  - c. ☒ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
  - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
  - b. ☐ have been transmitted by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
  - d. ☐ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

**Items 11. to 16. below concern document(s) or information included:**

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.  
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☐ Other items or information:

097/869251

17. ☒ The following fees are submitted:

**BASIC NATIONAL FEE ( 37 CFR 1.492 (a) (1) - (5) ) :**

Neither international preliminary examination fee (37 CFR 1.482)  
nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO  
and International Search Report not prepared by the EPO or JPO ..... \$970.00

International preliminary examination fee (37 CFR 1.482) not paid to  
USPTO but International Search Report prepared by the EPO or JPO. .... \$840.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but  
international search fee (37 CFR 1.445(a)(2)) paid to USPTO ..... \$690.00

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
but all claims did not satisfy provisions of PCT Article 33(1)-(4) ..... \$670.00

International preliminary examination fee paid to USPTO (37 CFR 1.482)  
and all claims satisfied provisions of PCT Article 33(1)-(4) ..... \$96.00

**ENTER APPROPRIATE BASIC FEE AMOUNT =**

**CALCULATIONS** PTO USE ONLY

\$ 690.00

Surcharge of \$130.00 for furnishing the oath or declaration later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(e)).

\$ 0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	27 - 20 =	7	X \$18.00
Independent claims	1 - 3 =	0	X \$78.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$260.00

\$ 126.00

\$ 0.00

\$ 0.00

**TOTAL OF ABOVE CALCULATIONS =**

\$ 816.00

Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement  
must also be filed (Note 37 CFR 1.9, 1.27, 1.28).

\$ 0.00

**SUBTOTAL =**

\$ 816.00

Processing fee of \$130.00 for furnishing the English translation later than ☐ 20 ☐ 30  
months from the earliest claimed priority date (37 CFR 1.492(f)).

\$ 0.00

**TOTAL NATIONAL FEE =**

\$ 816.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be  
accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property

\$ 80.00

**TOTAL FEES ENCLOSED =**

\$ 896.00

Amount to be  
refunded: \$  
charged: \$

a. ☐ A check in the amount of \$\_\_\_\_\_ to cover the above fees is enclosed.

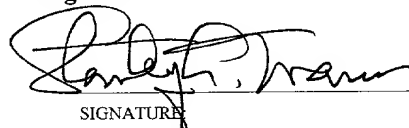
b. ☒ Please charge my Deposit Account No. 01-2717 in the amount of \$ 896.00 to cover the above fees.  
A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any  
overpayment to Deposit Account No. 01-2717. A duplicate copy of this sheet is enclosed.

**NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.**

SEND ALL CORRESPONDENCE TO

Stanley A. Marcus  
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2000 Market Street  
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US



SIGNATURE

Stanley A. Marcus

NAME

24,425

REGISTRATION NUMBER

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TACK FREE SURFACE CURES OF POLYMERS BY  
ORGANIC PEROXIDES IN THE PRESENCE OF AIR

This application claims priority from provisional application 60/164,488  
filed November 9, 1999.

BACKGROUND OF THE INVENTION

This invention relates to compositions of matter classified in the art of  
chemistry as bis-, tri- and higher poly-maleimides, as bis-, tri- and higher poly-  
citraconimides, as silicone elastomers, as p-phenylenediamine based antiozonants  
and as sulfur containing organic compounds which are accelerators for the sulfur  
curing (crosslinking) of polymers which are curable/crosslinkable by sulfur and  
also sulfur compounds which are polysulfide polymers. The invention also  
relates to compositions containing them, to processes for their use and to the  
products produced by such processes.

Polymers and copolymers crosslinked with free radical initiators, organic  
peroxides and/or azo initiators, are known to have superior properties,  
particularly to polymers crosslinked by sulfur cure. These properties include high

heat ageing resistance, low compression set, decreased staining of metal or coated metal sheet and easy production of colored products which have color stability during crosslinking and during long periods of use. These properties make use of peroxide cure of great practical importance particularly because crosslinking is through a carbon carbon bond rather than through a sulfur containing linkage and this bonding difference is responsible for the improved heat aging and compression set. The drawback for cure of polymers with free radicals from organic peroxides and azo initiators has always been that if air is not excluded from the surface of the material during cure, a tacky surface due to cure inhibition by the molecular oxygen in the air results.

In order to avoid tacky surfaces on objects fabricated using such free radical crosslinking by organic peroxides and/or azo initiators, it has been conventional to exclude air from contact with the surface during cure to avoid the cure inhibition caused by the presence of the molecular oxygen found in atmospheric air. Measures to exclude molecular oxygen add to the cost and complexity of the cure step and sometimes it is difficult, as in the cases of cure in steam autoclaves and in the interior of hoses, to assure the complete exhaustion of air and its molecular oxygen component. In some cases the manufacturer would like to switch from sulfur to peroxide cure and use existing hot air oven curing chambers. Curing with conventional peroxide systems under these circumstances would not be viable as a tacky surface would result.

In order to simplify and reduce the cost and complexity of the cure step, various methods have been suggested for preventing surface cure inhibition by molecular oxygen during free radical crosslinking. These methods have, for various reasons, met with little or no success in actual practice. In particular, none have provided a tack free surface while providing the most desirable physical property of peroxide (azo) cure; superior compression set at 150°C for 70 hours, compared to about 100°C, i.e. lower temperature performance for the prior art.

#### PRIOR ART

U. S. Patent 4,983,685 discloses the use of compounds selected from the

following classes: (a) imidazole compounds, (b) thiourea compounds, (c) thiazole compounds, (d) thiuram compounds, (e) dithiocarbamate compounds, (f) phenol compounds, (g) triazole compounds and (h) amine compounds which are accelerators for sulfur vulcanization in the optional presence of antioxidants, anti-ageing compounds and the like for elastomers for reducing surface tack in the peroxide cure of elastomers in the presence of molecular oxygen. Among optional ingredients which are suggested as possible ingredients for inclusion in the formulations, in this case for increased crosslinking, is N,N'-m-phenylene bismaleimide.

This is not the preferred optional coagent as a dimethacrylate compound is actually used in the examples. There is no recognition that this latter bismaleimide compound might provide an enhanced effect on the ability of certain compounds of (a) through (h) to reduce surface tack during free radical cure in the presence of molecular oxygen. The use of the various sulfur accelerators, in particular with peroxide, does provide tack free or reduced tack surfaces for the cured polymers in this reference but the important physical properties expected from a peroxide cure are also reduced. There is no recognition in U.S. Patent 4,983,685 that if the silicone elastomers, bismaleimides and biscitraconimides of the present invention are used in combination with p-phenylene diamine based antiozonants, sulfur containing sulfur vulcanization accelerators and antioxidants and/or polysulfide polymers in free radical cures of polymers, that tack free surfaces and improved physical properties will result and that of all the crosslinking aids mentioned, only these particular classes of compounds have that effect.

Japanese Published Patent Application No. Hei 9[1997]-169873 discloses that antioxidants of the benzimidazole type and of the polymeric 2,2,4-trimethyl-1,2-dihydroquinoline type used in combination with standard crosslinking aids such as methacrylate esters, triallylcyanurates and maleimides, such as this present invention's preferred component N,N'-m-phenylene bismaleimide, and standard crosslinking peroxides will result in cured peroxide crosslinkable elastomers with a tack free surface in the presence of air. The inclusion of p-

phenylene diamine based antiozonants, silicone elastomers, sulfur accelerators of any type and/or polysulfide polymers is not suggested.

U.S. Patent 4,334,043 teaches the use of surface treatment of curable polymer compositions with organo-metallic compounds, inorganic metallic salts, or lanthanides prior to crosslinking with organic peroxide crosslinking initiators in air to prevent surface tack after crosslinking. Other means of controlling surface tack are not mentioned except for the previously known techniques for surface tack free curing by simply excluding air contact with the rubber surface.

U.S. Patent 4,814,384 and 4,973,627 disclose cures of rubber blends for tire treads and sidewalls using a combination of sulfur and peroxide cures. Sulfur accelerators are also employed. Coagents of any type are not mentioned, nor is cure in the presence of air discussed. The use of elemental sulfur is required in the practice of these inventions. We have found, however, that the use of elemental sulfur adversely affects the final physical properties of the cured elastomer to the point where they are more typical of sulfur cure than peroxide cure.

U.S. Patent 4,743,656 also discloses a mixed sulfur/peroxide cure agent for elastomers, which cure agent also includes sulfur accelerators as well as elemental sulfur and the peroxide. Coagents are not mentioned, nor are crosslinking in air and surface tackiness discussed.

U.S. Patent 4,575,552 claims the use of specific combinations of hindered phenol antioxidants, metal salts of dithiocarbamates and m-phenylene-dimaleimide to provide a peroxide crosslinked polymer with superior hydrolytic and thermal stability for geothermal applications. There is no mention of crosslinking in the presence of air, air-inhibition or surface tackiness as a result of air-inhibition.

U.S. Patent 5,849,214 discloses the use of sulfur compounds, sulfur accelerators and hydroquinones with the optional presence of crosslinking aids (coagents) in the retardation of scorch during compounding of free radical crosslinkable polymers in the presence of free radical initiators. Bismaleimides, and biscitraconimides are not specifically discussed nor is there any mention of

the possible effect on surface tackiness during cure in the presence of molecular oxygen (air) for any of the compositions disclosed.

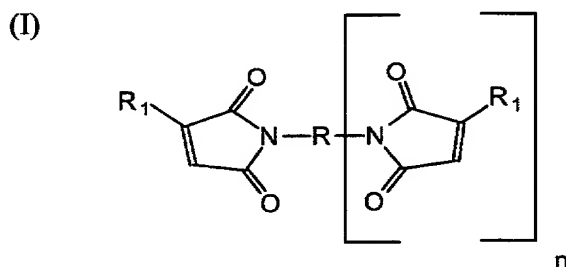
In addition there are patents which use various compounds to physically coat the surface of crosslinkable elastomers to exclude air (oxygen) e.g., US 4,439,388 which teaches use of boric acid, boric acid anhydride as a surface treatment prior hot air cure. This surface coating technique is labor intensive as it must be removed and disposed of after the crosslinking reaction step is completed.

None of the above references, taken singly or in combination, suggests applicants' solution described and claimed herein for elimination of surface tack due to air-inhibition during free radical cure of polymers by free radical curing agents, such as organic peroxides and azo initiators while providing the desirable physical properties of a peroxide (azo) cure. For example, the compression set values expected from a standard peroxide cure and wherein the compression set is measured at temperatures in the region about 150°C for 70 hours.

### SUMMARY OF THE INVENTION

The invention provides in its first composition aspect, a composition comprising:

- a) At least one compound (A) selected from the group consisting of silicone elastomers and a compound having the formula (I):



wherein n is 1, or 2 and R is divalent, or trivalent and is selected from the

group consisting of acyclic aliphatic groups having from about 2 to 16 carbon atoms, cyclic aliphatic groups having from about 5 to 20 carbon atoms, aromatic groups having from about 6 to 18 carbon atoms and alkyl aromatic groups having from about 7 to 24 carbon atoms, and wherein those divalent, or trivalent groups may contain one or more heteroatoms selected from O, N and S, replacing a carbon atom, or atoms, and each  $R_1$  is identical and is hydrogen or an alkyl group of 1 to 18 carbon atoms; and

(b) at least one compound (B) selected from the group consisting of p-phenylenediamine based antiozonants and sulfur containing organic compounds selected from the group consisting of sulfur containing organic compounds capable of accelerating sulfur vulcanization of polymers capable of being crosslinked by sulfur ("sulfur accelerators"), polysulfide polymers and mixtures of said sulfur containing compounds.

The tangible embodiments of the first composition aspect of the invention possess the inherent applied use characteristic of being suppressors of surface inhibition of free radical induced cure of polymers in the presence of gaseous molecular oxygen, (i.e., oxygen present in the atmosphere) thereby permitting tack free cures of polymers by free radical curing agents in the presence of air while maintaining the final physical properties associated with a conventional peroxide cure.

The invention provides in a subgeneric aspect of the first composition aspect of the invention, a composition formed by mixing as the essential ingredients thereof at least one member of compound (A) and at least one member of compound (B) of the first composition aspect of the invention.

The invention provides in a second composition aspect, a composition comprising a composition as defined in the first composition aspect and a free radical initiator selected from the group consisting of organic peroxides and azo initiators.

The tangible embodiments of the second composition aspect of the invention possess the inherent applied use characteristic of being curing or crosslinking agents for those polymers capable of being crosslinked by free radical initiators and of being capable of effecting such cures in the presence of



air (molecular oxygen) without the polymer being cured experiencing surface inhibition by the presence of air (molecular oxygen), thus, providing a cured or crosslinked polymer having a substantially tack free surface without the necessity for avoiding contact of said surface with air (molecular oxygen) during cure.

5           The invention provides a subgeneric aspect of the second composition aspect of the invention. Said composition being one prepared by mixing in any order, at least one member of compound (A) and at least one member of compound (B) of the first composition aspect of the invention and a free radical initiator as defined for the second composition aspect of the invention.

10           The invention provides in a third composition aspect a curable composition comprising a polymer curable by free radical initiators and a composition as defined in the second composition aspect of the invention.

15           The third composition aspect of the invention possesses the inherent applied use characteristic of being formable into a shaped article and then being crosslinkable while the surface of said shaped article is in contact with air (molecular oxygen) to provide a crosslinked shaped article having a substantially tack free surface.

20           The invention provides in a subgeneric composition aspect of the third composition aspect of the invention a curable composition prepared by mixing in any order at least one member of compound (A) and at least one member of compound (B) of the first composition aspect of the invention, a free radical initiator as defined for the second composition aspect of the invention and a polymer crosslinkable by a free radical initiator.

25           The invention provides in a first process aspect a process for the preparation of the second composition aspect of the invention which comprises mixing in any order at least one member of compound (A) and at least one member of compound (B) of the first composition aspect of the invention and a free radical initiator as defined in the second composition aspect of the invention.

30           The invention provides in a second process aspect of the invention, a process for the preparation of the third composition aspect of the invention which comprises mixing in any order at least one member of compound (A) and at least one member of compound (B) of the first composition aspect of the invention, a

free radical initiator as defined in the second composition aspect of the invention and a polymer crosslinkable by a free radical initiator.

Special mention is made of embodiments of the several aspects of the invention wherein compound (A) is selected from bismaleimides and compound (B) is selected from sulfur accelerators, where compound (A) is selected from biscitraconimides and compound (B) is selected from sulfur accelerators, where compound (A) is selected from bismaleimides and compound (B) is selected from polysulfide polymers, where compound (A) is selected from biscitraconimides and compound (B) is selected from polysulfide polymers, where compound (A) is selected from silicone elastomers and compound (B) is selected from polysulfide polymers.

Special mention is also made of embodiments of the several aspects of the invention wherein chlorinated polyethylene and/or chlorosulfonated polyethylene are included as optional supplemental ingredients in addition to compounds (A) and (B).

#### DETAILED DESCRIPTION

The best mode contemplated by the inventors for making and using their invention will now be described in detail with reference to a particular embodiment thereof, namely:

A mixture of dipentamethylene thiuram tetra-sulfide (Sulfads), N,N'-m-phenylene bismaleimide (HVA-2) and 1,1-di(t-butylperoxy)-3,3,5-trimethylcyclohexane (LUPEROX® 231 XL) used to cure ethylene propylene copolymer (VISTALON® 504) in hot air.

To prepare the mixture of Sulfads, HVA-2 and LUPEROX 231 XL, the ingredients, which are all in dry powder form (the LUPEROX® 231 XL is in the form of 40% by weight peroxide dispersed on calcium carbonate), may be mixed in any order and then compounded by standard methods (Banbury, two roll mill, extruder and the like) into the VISTALON® polymer. The Sulfads®, HVA-2 and LUPEROX 231 XL may also be compounded directly into the VISTALON either

simultaneously or sequentially in any order. Any two of the Sulfads, HVA-2 and LUPEROX 231 XL ingredients may be mixed and compounded into the VISTALON separately or simultaneously with the third ingredient. This compounding, if done separately, may also be performed in any order of ingredient addition to the polymer, but it is preferred if the peroxide is added last.

Once compounding with the VISTALON is complete, the compounded mixture may be cured simply by placing it in a hot air oven at a suitable temperature for initiating cure by decomposition of the peroxide, conveniently, in this case, at about 365°F (about 185°C), for a sufficient period of time to permit the desired degree of crosslinking to take place, conveniently, in this case, about 10 minutes, for a thin sample at room temperature at the start.

One of skill in the art will recognize that the other compounds falling within the scope of Formula I of the first composition aspect of this invention are all solid materials, are all trimaleimides, bismaleimides, tricitraconimides, or bis-citraconimides and can all be combined with the other starting materials contemplated by the invention by similar conventional methods to those described. The bismaleimides and biscitraconimides contemplated as starting materials are all either commercially available or can be readily synthesized by methods well known in the art. See, for example, U.S. Patents 5,484,948; 5,616,666, 5,292,815 and the references cited therein for more general synthetic methods.

The trimaleimides and tricitraconimides as well as the higher polymaleimides and citraconimides may be prepared by analogous techniques if they are not commercially available. For example, the trimaleimide, N,N',N''-(1,3,5-triazine-2,4,6-triyl)trimaleimide has CAS number CAS(67460-81-5).

Some primary amines suitable for synthesis of the di, tri- and higher polymaleimides and analogous citraconimides are polyfunctional primary amines such as melamine and the various polyoxypropylene amines such as the polyoxypropylene diamines and the polyoxypropylene triamines sold under the JEFFAMINE tradename by Huntsman Corporation.

In addition to the N,N'-m-phenylene-bismaleimide specifically referenced above, other bismaleimides, in addition to those disclosed in the above referenced

patents, suitable for use in the invention, without limiting the generality of the above general formula (I), are:

N,N'-ethylenebismaleimide, N,N'-hexamethylenebismaleimide, N,N'-dodecamethylene-bismaleimide, N,N'-(2,2,4-trimethylhexamethylene)  
 5 bismaleimide, N,N'-(oxy-dipropylene)bismaleimide, N,N'-(aminodipropylene)bismaleimide, N,N'-(ethylenedioxy-dipropylene)bismaleimide, N,N'-(1,4-cyclohexylene)bismaleimide, N,N'-(1,3-cyclohexylene) bismaleimide, N,N'-(methylene 1,4-dicyclohexylene) bismaleimide, N,N'-(isopropylidene-1,4-dicyclohexylene)bismaleimide, N,N'-(oxy-1,4-dicyclohexylene)bismaleimide, N,N'-p-(phenylene) bismaleimide, N,N'-(o-phenylene)bismaleimide, N,N'-(1,3-naphthylene)bismaleimide, N,N'-(1,4-naphthylene) bismaleimide, N,N'-(1,5-naphthylene)bismaleimide, N,N'-(3,3'-dimethyl-4,4'-diphenylene)bismaleimide, N,N'-(3,3-dichloro-4,4'-biphenylene)bismaleimide, N,N'-(2,4-pyridyl)bismaleimide, N,N'-2,6-  
 15 pyridyl)bismaleimide, N,N'-(1,4-anthraquinonediy)l bismaleimide, N,N'-(m-tolylene)bismaleimide, N,N'-(p-tolylene)bismaleimide, N,N'-(4,6-dimethyl-1,3-phenylene)bismaleimide, N,N'-(2,3-dimethyl-1,4-phenylene)bismaleimide, N,N'-(4,6-dichloro-1,3-phenylene)bismaleimide, N,N'-(5-chloro-1,3-phenylene) bismaleimide, N,N'-(5-hydroxy-1,3-phenylene) bismaleimide, N,N'-(5-methoxy-  
 20 1,3-phenylene) bismaleimide, N,N'-(m-xylylene)bismaleimide, N,N'-(p-xylylene)bismaleimide, N,N'-(methylenedi-p-phenylene) bismaleimide, N,N'-(isopropylidenedi-p-phenylene) bismaleimide, N,N'-(oxydi-p-phenylene)bismaleimide, N,N'-(thiodi-p-phenylene)bismaleimide, N,N'-(dithiodi-p-phenylene)bismaleimide, N,N'-(sulfodi-p-phenylene) bismaleimide,  
 25 N,N'-(carbonyldi-p-phenylene) bismaleimide,  $\alpha,\alpha$ -bis-(4-maleimodophenyl)-meta-diisopropylbenzene,  $\alpha,\alpha$ -bis-(4-p-phenylene) bismaleimide and  $\alpha,\alpha$ -bis-(4-maleimidophenyl)-para-diisopropylbenzene.

Combination of two or more bismaleimides, or bismaleimides with the trimaleimides, and with the higher polymaleimides in the compositions and  
 30 processes of the invention are also contemplated as equivalents and one of skill in the art would understand that such tri and higher polymaleimides and their substitution for the compounds and processes specifically illustrated herein for

the practice of the invention to be such equivalents and to be well within the scope contemplated by the invention.

Biscitraconimides, which may be substituted in whole or in part for the N,N'-m-phenylenebismaleimide referenced above include as representative examples:

- 1,2-N,N'-dimethylene biscitraconimide;
- 1,2-N,N'-trimethylene biscitraconimide;
- 1,5-N,N'-(2-methyl-pentamethylene)-biscitraconimide; and
- N,N'-methylphenylene biscitraconimide.

Mixtures of biscitraconimides and mixtures of bismaleimides and biscitraconimides as well as those including the trimaleimides are also contemplated as equivalents by the invention.

The biscitraconimides contemplated by the invention are all well known compounds and where not commercially available, they may be readily synthesized by methods detailed in the art. U.S. Patent 5,292,815 in column 4, provides a detailed list of such methods. As stated above, the tri- and higher polycitraconimides may be prepared by analogous methods and substituted in whole or in part in the compositions of the invention and such compounds and substitutions will be understood by one of skill in the art as being a full equivalent to those specifically illustrated herein and well within the scope contemplated as equivalent by the invention.

The silicone elastomers contemplated as useful in the aspects of the invention are the peroxide crosslinkable dimethyl vinyl substituted silicone derivative elastomers which are well known in the art. See, for example, "Kirk Othmer Encyclopedia of Chemical Technology", Vol. 20, pp. 943 et seq., John Wiley & Sons, © 1982.

Sulfur containing organic compounds capable of accelerating sulfur vulcanization of polymers, which are capable of being crosslinked by sulfur contemplated for use in the invention are well known in the art. Many different classes of these compounds are known and all are contemplated as equivalent.

*The Vanderbilt Rubber Handbook*, thirteenth edition, 1990, R.T. Vanderbilt Company, Inc., publisher lists many types. Illustrative of these are

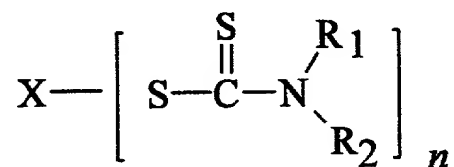
derivatives of benzothiazoles, thiadiazoles, sulfenamides, sulfenimides, dithiocarbamates, thiurams, imidazoles, xanthates, and thioureas. Also included in this general class of sulfur compound sulfur accelerators are sulfides, disulfides (e.g., diallyldisulfide) polysulfides and arylpolysulfide compounds such as the

5 amylphenol polysulfides e.g. VULTAC® products from ATOFINA Chemicals, Inc. and other sulfides such as disulfide and/or other known sulfur accelerating polysulfide phosphate, dithiophosphates and/or phosphorous and sulfur containing compounds. Other sulfur containing organic compounds capable of sulfur donation at vulcanization temperatures which are known but are not

10 presently used for such reactions because of cost concerns are also contemplated as equivalents. Illustrative of these is the compound 2-(2,4-cyclopentadiene-1-ylidene)-1,3-dithiolane.

More particularly, one sulfur accelerator class suitable for use in the practice of the invention are salts of disubstituted dithiocarbamic acid.

15 These salts have the general structure:



Wherein X is an ion derived from a metal selected from the group consisting of nickel, cobalt, iron, chromium, tin, zinc, copper, lead, bismuth, cadmium, selenium and tellurium, or X is a quaternary ammonium ion, *n* may vary

20 from 1 to 6 and is equal to the number of formal positive charges on the X ion, and *R*<sub>1</sub> and *R*<sub>2</sub> are independently alkyl of 1 to 7 carbon atoms.

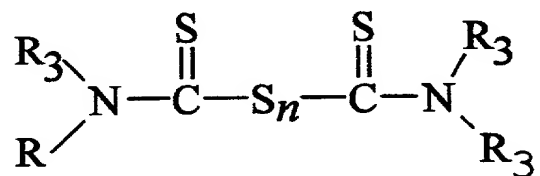
Examples of the salts of disubstituted dithiocarbamic acid are:

bismuth dimethyldithiocarbamate;

cadmium diethyldithiocarbamate;

cadmium diamyldithiocarbamate;  
 copper dimethyldithiocarbamate;  
 lead diamyldithiocarbamate;  
 lead dimethyldithiocarbamate;  
 5 selenium diethyldithiocarbamate;  
 selenium dimethyldithiocarbamate;  
 tellurium diethyldithiocarbamate;  
 piperidinium pentamethylene dithiocarbamate;  
 zinc diamyldithiocarbamate;  
 10 zinc diisobutyldithiocarbamate  
 zinc diethyldithiocarbamate;  
 zinc dimethyldithiocarbamate;  
 copper dibutyldithiocarbamate;  
 sodium dimethyldithiocarbamate;  
 15 sodium diethyldithiocarbamate;  
 sodium dibutyldithiocarbamate;  
 zinc di-n-butyldithiocarbamate;  
 zinc dibenzylthiocarbamate.

A second sulfur accelerator class suitable for use in the invention  
 20 comprises the thiurams. These are prepared from secondary amines and carbon  
 disulfide and possess the general structure:



25 Wherein R<sub>3</sub> is an alkyl group of from 1 to about 7 carbon atoms or the R<sub>3</sub>  
 groups on each particular nitrogen atom may be concatenated to form, together  
 with the nitrogen atom on which they are attached, a five, six or seven membered  
 heterocyclic ring containing 4, 5 or 6 carbon atoms respectively and *n* may have a

positive value from greater than zero up to 6.

Typical examples of thiuram sulfur accelerators are:

dipentamethylenethiuram tetrasulfide and hexasulfide;

tetrabutylthiuram disulfide;

tetramethylthiuram disulfide;

tetraethylthiuram disulfide;

tetramethylthiuram monosulfide;

isobutylthiuram disulfide;

dibenzylthiuram disulfide;

tetrabenzylthiuram disulfide;

tetraisobutylthiuram disulfide;

isobutylthiuram monosulfide;

dibenzylthiuram monosulfide;

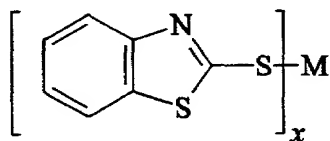
tetrabenzylthiuram monosulfide;

tetraisobutylthiuram monosulfide.

The higher multisulfides of the various thiurams are also sulfur donors.

Derivatives of thiadiazoles are, but not limited to, monobenzoyl derivatives of dimercaptothiadiazole (2,5-dimethyl-1.3.4-thiadiazole); the proprietary thiadiazole of the Vanderbilt Rubber Company identified as VANAX® 189; 1,2,4-thiadiazole, 5-ethoxy-3-(trichloromethyl) thiadiazole; and alkyl mercaptothiadiazoles, e.g. methyl mercapto thiadiazole.

Derivatives of benzothiazoles have the general structure:



Wherein M is a direct bond between two sulfur atoms, H, or an ion derived from a metal selected from the group consisting of nickel, cobalt, iron, chromium, tin, zinc, copper, lead, bismuth, cadmium, selenium and tellurium; and when M is H, x is 1; when M is a direct bond between two sulfur atoms, x is 1 or 2; and when



M is an ion derived from a metal,  $x$  is equal to the formal valence of the metal ion; and if M is a direct bond between two sulfur atoms and  $x$  is 1, then the second sulfur atom to which the M bond is attached is also bonded to a 4-morpholinyl radical.

5 Illustrative compounds are: 2-(4-morpholinodithio) benzothiazole; benzothiazyl disulfide; 2-mercapto-benzothiazole; 2-mercaptobenzothiazole disulfide; sodium-2-mercaptobenzothiazolate; zinc-2-mercapto-benzothiazole; copper-2-mercaptobenzothiazolate; 2-N-cyclohexylaminobenzothiazole; N-cyclohexylamino-2-benzothiazole polysulfide; 2-bisbenzothiazole-2,2'-polysulfide  
10 and 2-bisbenzothiazole-2, 2'-disulfide; bis(2,2'-benzothiazyl)disulfide).

The sulfenamide accelerators are also well known. Illustrative examples are: N-oxydiethylene-2-benzothiazole sulfenamide; N-oxydiethylene thiocarbonyl-N-oxydiethylene sulfenamide; N-cyclohexyl-2-benzothiazole sulfenamide; N-t-butyl-2-benzothiazole sulfenamide; N-cyclohexyl-2-  
15 benzothiazylsulfenamide; N,N-dicyclohexyl benzothiazyl sulfenamide; N-t-butyl-2-benzothiazole sulfenamide. There are also sulfenimide compounds, e.g., N-t-butyl-benzothiazole-2-sulfenimide.

Typical imidazoles are: 2-mercaptobenzimidazole, 2-mercaptomethylbenzimidazole; and the zinc salt of 2-mercaptobenzimidazole.

20 Zinc isopropyl xanthate is a typical xanthate sulfur accelerator.

Typical thioureas are: trimethylthiourea; 1,3-diethylthiourea and 1,3-dibutylthiourea; ethylene thiourea; blend of dialkyl thioureas; diphenyl thiourea; diorthotolyl thiourea; dimethyl thiourea; diethyl thiourea; dibutyl thiourea.

Alkylphenoldisulfide types of sulfur accelerators are illustrated by the  
25 compounds available from ATOFINA Chemicals, Inc., under the designation VULTAC® 2, VULTAC 3 and VULTAC 5.

Thiophosphate sulfur accelerators are illustrated by such compounds as copper dialkyldithiophosphate; zinc dialkyldithiophosphate; zinc amine dithiophosphate; zinc dibutyldithiophosphate; copper O,O-diisopropyl-  
30 phosphorodithiolate; zinc O,O-diisopropylphosphorodithiolate.

Other miscellaneous sulfur accelerators include 4,4-dithiodimorpholine; N,N'-caprolactam disulfide; dibutylxanthogen disulfide.

The polymers which can be cured (crosslinked) in the presence of molecular oxygen include all those natural and synthetic polymers capable of being crosslinked either by abstraction of hydrogen (or other extractable atoms, such as with iodo and bromo substituted fluoroelastomers) or by polymerization through double bonds.

Polymers which are currently understood to not be crosslinkable by these mechanisms and which undergo degradation in the presence of free radicals generated from organic peroxides and the azo initiators defined herein below and whose presence should be substantially avoided in the curable compositions of this invention include: poly(vinyl chloride), poly-(propylene), butyl rubber, epichlorohydrin polymers and epichlorohydrin ethylene oxide polymers. When used herein and in the appended claims when referring to polymers suitable for use in connection with the invention any reference to a group of polymers using the terms "comprising," "consisting essentially" and "consisting of" expressly excludes more than minor insignificant amounts (1% by weight or less) of the non free radical crosslinkable polymers in the absence of an express statement to the contrary.

Polymers crosslinkable by free radicals from organic peroxides and azo initiators as defined herein below include ethylene-propylene terpolymer (EPDM), ethylene-propylene copolymer (EPM) natural polyisoprene rubber (NR), styrene butadiene rubber (SBR), polybutadiene rubber (BR), synthetic polyisoprene rubber (IR), poly(ethylene) (PE), ethylene-vinyl acetate (EVA), acrylonitrile-butadiene-styrene (ABS), unsaturated polyester, styrene-butadiene-styrene block copolymers (SBS), styrene-isoprene-styrene block copolymers (SIS), neoprene rubber (CR), nitrile rubber (NBR), polysulfide rubber (T) chlorinated poly-(ethylene) (CM), polyurethane (AU, EU), vinylidene fluoride copolymers (CFM), silicone rubber (PMQ), vinyl silicone rubber (VMQ, PVMQ), polyacrylate (ACM), chlorosulfonated poly(ethylene) (CSM) and fluorosilicone rubber (FVMQ).

The free radical initiators (organic peroxides and azo initiators) suitable for use in the invention include all those classes of organic peroxides and azo initiators suitable for curing (crosslinking) polymers, both thermoplastics and elastomers.

The azo initiators are those known in the art, such as 2,2'-azobis-(2-

acetoxyp propane), to generate free radicals on heat decomposition capable of inducing the desired curing (crosslinking) reaction. The azo initiators of U.S. Patents 3,862,107 and 4,129,531, the disclosures of which are incorporated herein by reference, are also suitable.

5 With the exception of hydroperoxides and liquid peroxydicarbonates, all those organic peroxides known to undergo decomposition by heat to generate radicals capable of initiating the desired curing (crosslinking) reactions are contemplated as suitable for use in the invention. Dialkyl peroxides, diperoxyketals, mono-peroxy carbonates, cyclic ketone peroxides, diacyl peroxides, 10 organosulfonyl peroxides, peroxyesters and solid, room temperature stable peroxydicarbonates are the preferred initiators. The most preferred initiators are dialkyl peroxides, peroxyketals, cyclic ketone peroxides and diacyl peroxides.

A good reference which provides important peroxide names and physical properties for all these classes of organic peroxides can be found in "Organic Peroxides" by Jose Sanchez and Terry N. Myers; Kirk-Othmer Encyclopedia of 15 Chemical Technology, Fourth Ed., Volume 18, (1996).

Illustrative dialkyl peroxide initiators are:

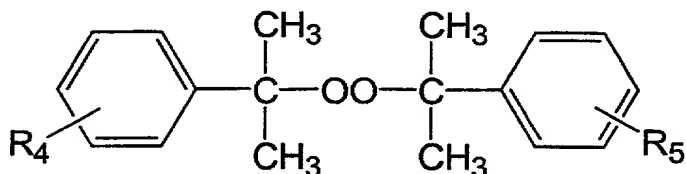
di-t-butyl peroxide;  
t-butyl cumyl peroxide;  
20 2,5-di(cumylperoxy)-2,5-dimethyl hexane;  
2,5-di(cumylperoxy)-2,5-dimethyl hexyne-3;  
4-methyl-4-(t-butylperoxy)-2-pentanol;  
4-methyl-4-(t-amylperoxy)-2-pentanol;  
4-methyl-4-(cumylperoxy)-2-pentanol;  
25 4-methyl-4-(t-butylperoxy)-2-pentanone;  
4-methyl-4-(t-amylperoxy)-2-pentanone;  
4-methyl-4-(cumylperoxy)-2-pentanone;  
2,5-dimethyl-2,5-di(t-butylperoxy)hexane;  
2,5-dimethyl-2,5-di(t-amylperoxy)hexane;  
30 2,5-dimethyl-2,5-di(t-butylperoxy)hexyne-3;  
2,5-dimethyl-2,5-di(t-amylperoxy)hexyne-3;  
2,5-dimethyl-2-t-butylperoxy-5-hydroperoxyhexane;

2,5-dimethyl-2-cumylperoxy-5-hydroperoxy hexane;  
 2,5-dimethyl-2-t-amylperoxy-5-hydroperoxyhexane;  
 m/p-alpha, alpha-di[(t-butylperoxy)-isopropyl]benzene;  
 1,3,5-tris(t-butylperoxyisopropyl) benzene;  
 1,3,5-tris(t-amylperoxyisopropyl) benzene;  
 1,3,5-tris(cumylperoxyisopropyl) benzene;  
 di[1,3-dimethyl-3-(t-butylperoxy)butyl]carbonate;  
 di[1,3-dimethyl-3-(t-amylperoxy)butyl]carbonate;  
 di[1,3-dimethyl-3-(cumylperoxy)butyl]carbonate;  
 di-t-amyl peroxide;  
 t-amyl cumyl peroxide;  
 2,4,6-tri (butylperoxy)-s-triazine;  
 1,3,5-tri [1-(t-butylperoxy)-1-methylethyl] benzene  
 1,3,5-tri-[(t-butylperoxy)-isopropyl]benzene;  
 1,3-dimethyl-3-(t-butylperoxy)butanol;  
 1,3-dimethyl-3-(t-amylperoxy)butanol; and mixtures thereof.

Illustrative solid, room temperature stable peroxydicarbonates are, but not limited to:

di(2-phenoxyethyl) peroxydicarbonate; di(4-t-butyl-cyclohexyl)  
 peroxydicarbonate; dimyristyl peroxydicarbonate; dibenzyl peroxydicarbonate;  
 di(isobornyl) peroxydicarbonate.

Another preferred class of dialkylperoxides which may be used singly or in  
 combination with the other free radical initiators contemplated by the invention are  
 those selected from the group represented by the formula:



Wherein R<sub>4</sub> and R<sub>5</sub> may independently be in the meta or para positions and  
 are the same or different and are selected from the group hydrogen or straight or  
 branched chain alkyl of from 1 to 6 carbon atoms. Dicumyl peroxide and

isopropylcumyl cumyl peroxide are illustrative.

Other dialkyl peroxides are:

3-cumylperoxy-1,3-dimethylbutyl methacrylate;

3-t-butylperoxy-1,3-dimethylbutyl methacrylate;

5 3-t-amylperoxy-1,3-dimethylbutyl methacrylate;

tri(1,3-dimethyl-3-t-butylperoxy butyloxy)vinyl silane;

1,3-dimethyl-3-(t-butylperoxy)butyl N-[1-{3-(1-methylethenyl)-  
phenyl}-1-methylethyl]carbamate;

10 1,3-dimethyl-3-(t-amylperoxy)butyl N-[1-{3-(1-methylethenyl)-  
phenyl}-1-methylethyl]carbamate;

1,3-dimethyl-3-(cumylperoxy)butyl N-[1-{3-(1-methylethenyl)-  
phenyl}-1-methylethyl]carbamate.

In the group of diperoxyketal initiators, the preferred  
initiators are:

15 1,1-di(t-butylperoxy)-3,3,5-trimethylcyclohexane;

1,1-di(t-butylperoxy)cyclohexane;

n-butyl 4,4-di(t-amylperoxy)valerate;

ethyl 3,3-di(t-butylperoxy)butyrate;

2,2-di(t-amylperoxy)propane;

20 3,6,6,9,9-pentamethyl-3-ethoxycarbonylmethyl-

1,2,4,5-tetraoxacyclononane;

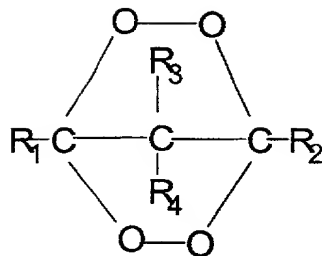
n-butyl-4,4-bis(t-butylperoxy)valerate;

ethyl-3,3-di(t-amylperoxy)butyrate; and mixtures thereof.

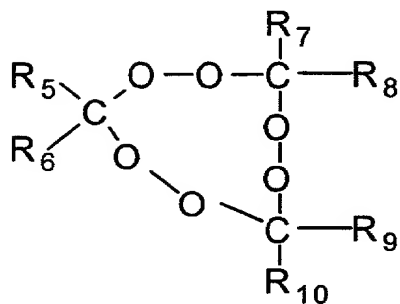
25 Other peroxides falling within the general class defined as useful in the  
invention include benzoyl peroxide, 00-t-butyl-0-hydrogen-monoperoxy-succinate  
and 00-t-amyl-0-hydrogen-monoperoxy-succinate.

Illustrative cyclic ketone peroxides are compounds having the general  
formulae (II), (III) and/or (IV).

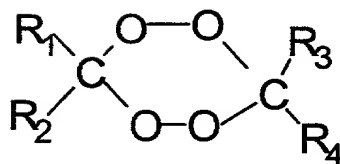
(II)



(III)



(IV)



- 5           Wherein  $R_1$  to  $R_{10}$  are independently selected from the group consisting of hydrogen,  $C_1$  to  $C_{20}$  alkyl,  $C_3$  to  $C_{20}$  cycloalkyl,  $C_6$  to  $C_{20}$  aryl,  $C_7$  to  $C_{20}$  aralkyl and  $C_7$  to  $C_{20}$  alkaryl, which groups may include linear or branched alkyl properties and each of  $R_1$  to  $R_{10}$  may be substituted with one or more groups selected from hydroxy,  $C_1$  to  $C_{20}$  alkoxy, linear or branched  $C_1$  to  $C_{20}$  alkyl,  $C_6$  to  $C_{20}$  aryloxy, halogen, ester, carboxy, nitride and amido, preferably, at least 20% of the total active oxygen
- 10           content of the peroxide mixture used for a crosslinking reaction will be from

compounds having formulas (II), (III) and/or (IV).

Some examples of suitable cyclic ketone peroxides are:

3,6,9, triethyl-3,6,9-trimethyl-1,4,7-triperoxynonane (or methyl ethyl ketone peroxide cyclic trimer) and methyl ethyl ketone peroxide cyclic dimer;

5 3,3,6,6,9,9-hexamethyl-1,2,4,5-tetraoxacyclononane;

Illustrative suitable peroxy esters are:

2,5-dimethyl-2,5-di(benzoylperoxy) hexane;

t-butylperbenzoate;

t-butylperoxy acetate;

10 t-butylperoxy-2-ethyl hexanoate;

t-amyl perbenzoate;

t-amyl peroxy acetate;

t-butyl peroxy isobutyrate;

3-hydroxy-1,1-dimethyl t-butyl peroxy-2-ethyl hexanoate;

15 00-t-amyl-0-hydrogen-monoperoxy succinate;

00-t-butyl-0-hydrogen-monoperoxy succinate;

di-t-butyl-diperoxyphthalate;

t-butylperoxy (3,3,5-trimethylhexanoate);

1,4-bis (t-butylperoxycarbo) cyclohexane;

20 t-butylperoxy-3,5,5-trimethylhexanoate;

t-butyl-peroxy-(cis-3-carboxy) propionate;

allyl 3-methyl-3-t-butylperoxy butyrate.

Illustrative monoperoxy carbonates are:

00-t-butyl-0-isopropylmonoperoxy carbonate;

25 00-t-butyl-0-(2-ethyl hexyl) monoperoxy carbonate;

1,1,1-tris[2-(t-butylperoxy-carbonyloxy)ethoxymethyl]propane;

1,1,1-tris[2-(t-amylperoxy-carbonyloxy)ethoxymethyl]propane;

1,1,1-tris[2-(cumylperoxy-carbonyloxy)ethoxymethyl]propane;

OO-t-amyl-O-isopropylmonoperoxy carbonate.

30 Illustrative diacyl peroxides are:

di(4-methylbenzoyl) peroxide;

di(3-methylbenzoyl) peroxide;  
di(2-methylbenzoyl) peroxide;  
didecanoyl peroxide; dilauroyl peroxide;  
2, 4-dibromo-benzoyl peroxide;  
5 succinic acid peroxide.  
dibenzoyl peroxide;  
di (2,4-dichloro-benzoyl) peroxide.

Imido peroxides of the type described in PCT Application publication  
WO9703961 A1 6 Feb 1997 are also contemplated as suitable for use by the  
10 invention.

One of skill in the art will readily be able to select suitable quantities of the  
various ingredients for use in the invention and will quickly and easily be able to  
optimize the concentrations through a series of bench scale trials employing  
increasing amounts of the ingredients in samples of the polymer to be cured  
15 (crosslinked). The optimum processing (compounding) time and temperatures and  
the like may also be determined from the same trials as will the optimum cure time  
and temperature.

Typically, one will employ the compounds of formula (a) (the bismaleimide  
and biscitraconimides) in the composition of the invention in quantities which will  
20 provide from about 0.2 parts by weight per part of polymer by weight (phr) to about  
10.0 phr, preferably from about 1.0 phr to about 5.0 phr, most preferably from about  
1.5 phr to about 3.0 phr.

Typically, one will employ the sulfur containing organic compound capable  
of accelerating sulfur vulcanization in polymers capable of being crosslinked by  
25 sulfur in compositions of the invention in quantities which will provide from about  
0.01 phr to about 20 phr, preferably from about 0.1 phr to about 1.0 phr, most  
preferably from about 0.1 phr to about 0.5 phr. It is understood by those of skill in  
the art that these compounds are of two types. Those that donate sulfur to the  
vulcanization and those which simply accelerate sulfur vulcanization. Either class of  
30 compound or mixtures thereof are contemplated as equivalents by the invention.

Alkyl phenol disulfide polymers of the VULTAC® type are preferably used  
at from about 0.5 phr to 20 phr when used alone or at from about 0.1 phr to about 10



phr when in combination with other sulfur accelerators.

Typically, one will employ the free radical initiator (organic peroxide and/or azo initiator) in quantities of from about 0.04 to about 10 phr preferably from about 1 to about 4 phr.

5       The time-temperature conditions necessary for curing largely depend on the structure of the free radical curing agent. For the azo initiators, suitable conditions are detailed in U.S. Patents 3,632,107 and 4,129,531.

10       For the compositions of the invention, appropriate time and temperature conditions may be determined for crosslinking a particular polymer composition by running a small number of well controlled rheometer studies and selecting values from the results of those studies where the time/temperature relationship is from five to fifteen times the half life value for the free radical initiator in the system.

15       The invention contemplates that other conventional additives such as anti-oxidants (hindered phenols and polymeric quinoline derivatives are preferred), aliphatic process oils, and other process aids, pigments, dyes, tackifiers, waxes, reinforcing aids, UV stabilization agents, blowing agents and activators and antiozonants may also be present in the compositions before, after and during the curing step.

20       The polysulfide polymers contemplated by the invention are those known polysulfide polymers which are prepared by the reaction of an  $\alpha,\omega$  - dihaloalkyl (or dihaloheteroalkyl) compound with a metallic, preferably an alkali metal, polysulfide. The common commercially available polysulfide polymers are either liquids or solids, are either thiol or hydroxy terminated and are derived from materials produced by the reaction of 1,2-dichloroethane, 2,2'-dichloro-diethyl ether or  
25       bis(2-chloroethyl)formal with an alkali metal polysulfide ( $MS_x$ ) wherein M is an alkali metal ion, preferably derived from sodium and x is a number greater than 1 up to about six.

30       The invention contemplates that polysulfide polymers may be used in place of or in admixture with the other compounds (B) in equal quantities to those previously specified for those compounds. Since an excess of polysulfide polymer is not contemplated as detrimental to the practice of the invention, it is also contemplated that they may be preblended with the compound(s) (A) and optionally with the free

radical initiator(s) to form masterbatches, either solid or liquid. The polysulfide polymers may also be preblended into the polymer to be cured and the compound(s) (A) and also the free radical initiator(s) blended in simultaneously or subsequently at the option of the operator. Use of the polysulfide polymers in combination with the other sulfur accelerators contemplated by the invention permits reduction of the amount of sulfur accelerator required by the invention.

Similarly it will be obvious to one of skill in the art that polysulfide polymers themselves may be cured to tack free surfaces with free radical initiators in the presence of molecular oxygen if a compound (A) is present in the curable composition even if another compound (B) is not present. Thus, the invention contemplates such a curable composition as an equivalent to the second composition aspect of the invention as defined above.

Certain crosslinkable elastomer compositions which are highly filled with oil and/or carbon black (commonly referred to as highly extended elastomer formulations) are normally cured using sulfur vulcanization rather than free radical initiators. Free radical cure is more difficult because the radicals generated lack specificity and react with the filler and oil as well as the elastomer. This reduces efficiency of the free radical initiator.

It has been found that the use of chlorinated polyethylene and /or chlorosulfonated polyethylene as supplemental ingredients to all the compositions of the various composition aspects of the invention surprisingly increases free radical cure efficiency in highly extended elastomer formulations and allows free radical cure of the systems with reduced or no surface tack. The amount of chlorinated and/or chlorosulfonated polyethylene as supplemental ingredients in the compositions of the first composition aspect of the invention may be from about 1% to about 50% by weight, preferably 15% to 40% by weight and more preferably from 20% to 35% by weight.

The inclusion of these two polymers as supplemental ingredients in some cases has been found to permit use of lower concentrations of the compositions of the first composition aspect of the invention in formulation of the compositions of the second composition aspect of the invention.

The following examples further illustrate the best mode contemplated by the

inventors for the practice of their invention and are to be construed as illustrative and not in limitation thereof.

### EXAMPLES

5     Example 1 - Evaluation of prior art compositions for crosslinking EPM in the presence of air and their effect on surface tack and level of crosslinking.

10     In the example, various known co-curing agents, e.g., unsaturated monomeric crosslinking coagents, elemental sulfur, or sulfur donor compounds, were evaluated in a cure with an organic peroxide. The elastomer used in this example was an ethylene propylene copolymer (EPM) marketed by Exxon, VISTALON® 504. Prior art (U.S. 4,983,685) claims that use of high levels (2.5 to 20 parts) of specific sulfur containing compounds with or without optional unsaturated monomers can be used to crosslink elastomers with peroxide in the presence of air and produce a non-tacky surface. Other references employ sulfur. In this example, low levels of sulfur or sulfur donor compounds were evaluated alone or with some monomeric coagents. 15     The data in Table 1 shows these formulations produced a sticky surface, but also provided a crosslinked product with good physical properties (higher MH torque). A blend (as taught by the prior art) of an optional monomeric coagent, a high level of sulfur compound and peroxide was also evaluated. This produced a non tacky surface when crosslinking elastomers in the presence of air, however, the physical 20     properties were unacceptable compared to a conventional peroxide monomeric crosslinking formulation.

25     In the formulation of Table 1, a standard carbon black formulation was used. Carbon black (N774), Sunpar® LW150 process oil from sun, zinc oxide, an antioxidant (AgeRite MA) were compounded into the EPM. In addition, all formulations included 8 parts of a 40% assay organic peroxyketal peroxide, LUPEROX® 231XL [40% 1,1-di(butylperoxy)-3,3,5-trimethylcyclohexane] dispersed on calcium carbonate, marketed by ATOFINA Chemicals, Inc. The level of crosslinking was determined using a Flexsys MDR® 2000E moving die rheometer. Samples were also cured in hot air at 365°F for ten minutes and judged

for surface tackiness by placing a paper towel on the surface, immediately after removal from the oven using moderate and consistent pressure. Surface tack was rated from 1-10, where 1 is considered non-tacky and 10 is very tacky.

A combination of peroxide and a low level of elemental sulfur, 0.3 parts (a typical EPM cure), (Table 1, run #1) provided a good level of crosslinking based on the  $M_H$  but a sticky surface when cured in hot air. Use of 2 parts HVA-2 (N,N'-phenylene bismaleimide, marketed by Dupont) coagent together with peroxide (Table 1, run #2) provides an expected improved level of crosslinking (based on  $M_H$ ), but the surface tackiness is, unfortunately, poor. Note: in each case, 1 part of an antioxidant, AgeRite MA (a polymerized quinoline marketed by R.T. Vanderbilt), is used, a standard level which does not affect the surface. A combination of 0.3 parts elemental sulfur and 2 parts of HVA-2 (Table 1, run #5) provides further improvement in the level of crosslinking, (compared to run #2 using HVA-2 alone), but, unfortunately, provides high surface tackiness (7 out of 10). Thus, blends of coagent with low levels of elemental sulfur can improve crosslinking, but not surface tackiness. Prior art teaches 2.5 to 20 parts of sulfur accelerator or its equivalent is required. The use of the coagent HVA-2 and peroxide in the carbon black EPM formulation containing oil and antioxidant provided good crosslinking, but, unfortunately, also produced a sticky surface when curing in hot air. Using 4 parts of monomeric coagent SR-350® together with 0.3 parts of elemental sulfur provided the highest level of crosslinking, but poor surface tackiness (see Table 1, run #4). SR-350 is trimethylolpropanetrimethacrylate, marketed by Sartomer.

Table 1, run #3, 1.2 parts of Sulfads® in combination with peroxide, provided an unacceptable surface tackiness of 6 out of 10, and an undesirable level of cure based on  $M_H$ . Sulfads is 98% dipentamethylene thiuram tetrasulfide, marketed by R. T. Vanderbilt; thus, the amount of dipentamethylene thiuram tetrasulfide actually added was  $1.2 \times 0.98 = 1.18$  parts. Finally, using 1.2 parts of Sulfads in combination with 2 parts HVA-2 and peroxide results in a good surface when cured in the presence of air, unfortunately, the desired crosslinked physical properties are reduced, as shown by the low  $M_H$  value attained.

This example illustrates that one cannot predict the effect of blending dissimilar co-curing agents such as peroxide, monomeric coagents and elemental sulfur or sulfur containing compounds with the objective of maintaining an acceptable level of crosslinking together with low surface tackiness when curing in the presence of air. One may postulate that if the overall level of crosslinking can be improved with various coagents and peroxide, the surface tackiness in the presence of air can be improved. However, our data shows that one can improve the level of crosslinking while not improving the surface tackiness when curing in the presence of air.

In addition, when the surface tack was finally reduced using teaching from the prior art, it was found that the overall crosslinking was undesirable. Crosslinking within the sample (excluding air) and crosslinking the surface in the presence of air are two separate processes. Increasing or decreasing the level of crosslinking coagents or use of the various co-curing agents in combination with peroxide leads to unpredictable results. Crosslinking coagents such as HVA-2 help to increase the level of crosslinking with peroxides cures, but appear to provide a poor surface tack in air cure. When using monomeric coagents, e.g., HVA-2 or SR 350 in combination with other co-curing agents, such as elemental sulfur or sulfur donor/accelerator compounds, as taught in the art, we find that the final physical properties are severely reduced, thus negating the advantages of a monomeric coagent-peroxide cure system.

**TABLE 1**

Run #	1	2	3	4	5	6
VISTALON® 504 EPM elastomer	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.						
N 774 (phr) Carbon Black	75	75	75	75	75	75
LW150 Oil phr	10	10	10	10	10	10
ZnO (phr)	5	5	5	5	5	5
AgeRite MA (antioxidant)	1	1	1	1	1	1
LUPEROX® 231XL 40% Peroxide	8	8	8	8	8	8
SR-350® (phr)	-	-	-	4.0	-	-
HVA-2	-	2.0	-	-	2.0	2.0
Sulfur	0.3	-	-	0.3	0.3	-
Sulfads® (98%)	-	-	1.2	-	-	1.2
Crosslinking, MDR 2000E Moving Die Rheometer at 150°C						
M <sub>H</sub> in. - lb.	29	39	20	44	43	29
Curing in hot air for 10 minutes at 365°F to determine surface tack						
Surface Tack*	7	7	6	10	7	1

\*Surface Tackiness Rating: 1 - 10 where 10 is very tacky and 1 is no tack.

Example 2 - Compounding of prior art peroxide formulations versus novel peroxide formulations for crosslinking fully saturated (no double bonds within the polymer chains) elastomers.

The novel peroxide formulation provides good physical properties associated with a conventional peroxide cure, plus an unexpected non-tacky surface when crosslinking in the presence of air.

In this example, 40% assay dicumyl peroxide dispersed on clay, was used to crosslink EPM (polyethylene propylene copolymer) VISTALON ® 707 (Exxon). Table 2, run #1, dicumyl peroxide blended with SR 206 (ethylene glycol dimethacrylate) was evaluated for crosslinking in the MDR, % compression set and in hot air cure for surface tack. Note: prior art (U.S. 4,983,685) used SR 206 as an optional monomeric coagent in their examples. This peroxide and coagent blend produced a good cure ( $M_H$ ) and % compression set, with a very sticky surface upon hot air cure at 400°F. Addition of Sulfads ® (98% dipentamethylene thiuram tetra sulfide) at the prior art claimed levels of 2.5 parts to the peroxide SR 206 blend improved the surface but totally destroyed % compression set; Table 2, run #2.

Using prior art levels in the specification of U.S. 4,983,685 of 0.8 parts of Sulfads, Table 2, run #4 together with an optional 2.2 parts HVA-2 coagent, 5 parts DC40 provided good surface, but very poor % compression set. Note: percent compression set values of 100% to 60% would be expected for sulfur vulcanization, not peroxide crosslinking.

Table 2, run #3, prior art taught levels of 5 parts Vanox ZMTI (93% Zinc 2-mercapto-toluidimidazole) with optional SR-206 coagent, improved compression set but gave a very sticky, unacceptable surface.

Unexpectedly, Table 2, run #5, low levels of dipentamethylene thiuram tetra sulfide (at 0.4 parts which falls outside of U.S. 4,983,685), 0.4 parts of Durax (98% N-cyclohexyl-2-benzothiazolesulfenamide) and 5 parts DC40 (40% dicumyl peroxide), provides a tack free surface when cured in a hot air oven with an improved 41% compression set and better level of crosslinking.

Table 2, run #6, shows that, unexpectedly, 0.4 parts Durax, 0.4 parts of Vanax A (98% 4,4'-dithiodimorpholine), 2.2 parts HVA-2 monomeric coagent and 5 parts DC40 (40% dicumyl peroxide), provided a tack free surface with 31%

compression set. The use of the Durax and/or Vanax A, with peroxide and bismaleimide type coagent, for use in hot air cure of elastomers, has not been taught in the prior art. Table 2, run 7, use of 0.8 parts of Durax, instead of the Durax/Vanax blend, provided a further unexpected outstanding % compression set value of 21% with a non-tacky surface upon hot air cure at 400°F.

Lastly, Table 2 run #8, unexpectedly outstanding results were obtained; an excellent, tack free surface with very desirable, extremely low 18% compression set using the novel blend of 2.2 parts HVA-2, 0.4 parts VULTAC 5, 0.4 parts Durax and 5 parts DC40. VULTAC 5 (75% alkyl phenol disulfide polymer from Elf Atochem) has not been taught by the art for use in hot air cure of elastomers together with peroxide and bismaleimide type coagent. Note that 0.4 parts of VULTAC 5 is equivalent to adding 0.3 parts of the alkyl phenol disulfide polymer, due to the 75% assay.

This example shows that prior art formulations using peroxide and high levels of select compounds, such as dipentamethylene thiuram tetra sulfide, blended with optional monomeric coagents do, in fact, produce a tack-free surface but provide very poor final physical properties for the crosslinked elastomer. Using 0.8 parts of a thiuram preferred compound taught by the prior art's U.S. 4,983,685 disclosure, but less than the 2.5 to 20 part levels claimed, produced a crosslinked elastomer with very poor heat aging properties for a peroxide cure formulation.

The high 60%+ compression set values show that the prior art crosslinked polymer permanently deformed under applied pressure over 70 hours at 150°C, a standard test (ASTM D-395-61) for a peroxide cure. A tack free surface can be obtained in hot air by sulfur vulcanization for unsaturated polymers, but results in poor heat aging properties. There is no advantage to a peroxide cure system cured in hot air over a sulfur vulcanization if there are no significant differences in physical properties, e.g., low  $M_H$  and/or high (poor) percent compression set values. Peroxide cures have been traditionally the choice for a number of high temperature and high performance applications. Peroxide crosslinking provides carbon-carbon bonds which allow the crosslinked polymer manufacturer to use the full engineering capabilities of the elastomer.



**TABLE 2**

Run #	1	2	3	4	5	6	7	8
VISTALON 707 EPM	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.								
N774 Carbon Black	75	75	75	75	75	75	75	75
LW150 process oil	30	30	30	30	30	30	30	30
ZnO	5	5	5	5	5	5	5	5
Stearic acid	1	1	1	1	1	1	1	1
AgeRite MA	1	1	1	1	1	1	1	1
DC40	8.0	8.0	8.0	5.0	5.0	5.0	5.0	5.0
SR-206	2.0	2.0	2.0	-	-	-	-	-
HVA-2	-	-	-	2.2	2.2	2.2	2.2	2.2
Sulfads (98%)	-	2.5	-	0.8	0.4	-	-	-
ZMTI (93%)	-	-	5.0	-	-	-	-	-
Vanax A (98%)	-	-	-	-	-	0.4	-	-
VULTAC 5 (75%)	-	-	-	-	-	-	-	0.4
Durax (98%)	-	-	-	-	0.4	0.4	0.8	0.4
MDR Cure at 170°C								
M <sub>H</sub> in-lb	17	9	16	17	18	17	16	16
T <sub>S1</sub> min	1.04	1.56	1.06	1.20	1.11	0.61	0.58	0.63
T <sub>C90</sub> min	5.8	7.5	5.5	6.6	5.2	5.9	4.1	4.2
Cure in 400°F air for 12 min.								
Surface	10	1.5	7	1.5	2	2.5	2.5	2
Compression Set 70 hr. at 150°C								
% Compression Set	25	100	35	63	41	31	21	18

Example 3 - Novel peroxide-additive compositions produce solid or foamed crosslinked elastomer in hot air, with outstanding physical properties, e.g., percent compression set, while also producing a non tacky surface when curing in the presence of air.

5           The elastomer used in this example was Uniroyal ® X3378 EPDM containing 4% dicyclopentadiene as the termonomer. Peroximon DC 40KEP (40% dicumyl peroxide on clay) was used as the peroxide crosslinking agent. In this example the levels of Sulfads® (98% dipentamethylene thiuram tetra sulfide), VULTAC ®5 (75% alkyl phenol disulfide polymer) and Durax® (98% N-  
10       cyclohexyl-2-benzothiazolesulfenamide) were varied, keeping the peroxide and HVA-2 coagent constant (Table 3, runs #2 to #6).

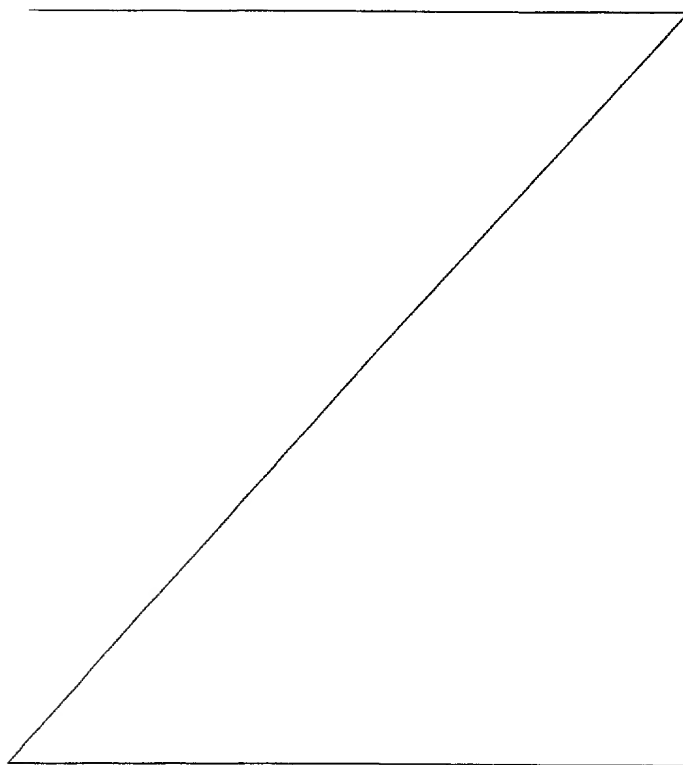
Note: adding 0.5 parts (phr) Sulfads is equivalent to 0.49 parts of dipentamethylene thiuram tetra sulfide, per 100 parts of rubber. Referring to runs #2, #3 and #4, unexpectedly, excellent compression set and a tack free surface were  
15       found when using dipentamethylene thiuram tetra sulfide below 0.5 phr, on a pure basis with HVA-2 and peroxide.

          When the Sulfads (98% dipentamethylene thiuram tetra sulfide), VULTAC 5 and Durax concentration were varied, it was unexpectedly discovered that both solid and foamed crosslinked elastomers could be produced by hot air cure using  
20       our novel peroxide/additive composition. Foamed crosslinked elastomer with a tack free surface was obtained when using <0.5 phr pure dipentamethylene thiuram tetra sulfide, with either VULTAC 5 or Durax; see Table 3, runs #2, #3 and #4. Using VULTAC 5 or a blend of VULTAC 5 with Durax provided solid crosslinked EPDM, Table 3, runs #5 and #6. Unexpectedly in each case (Table 2, runs #2 to  
25       #6) outstanding percent compression set values were obtained, virtually equivalent to a conventional peroxide cure, provided in Table 2, run #1.

          Sulfur vulcanization compression sets are measured at approximately 100°C - 120°C and rarely determined at 150°C due to constant poor performance at elevated temperature. The present 150°C, 70 hour desirable compression set data  
30       proves that these novel peroxide-additive compositions illustrated in the practice of this invention produce crosslinked EPDM with unexpectedly good heat aging properties. In addition, the novel peroxide-additive compositions surprisingly

provide a way to produce both solid and foamed articles.

- 5 Although not wishing to be bound, it is thought that faster scorch times related to the unexpected crosslinked network sets up faster than gas evolution from the cure system early in the cure to provide solid cured articles, which is more difficult under low pressure conditions in an oven as opposed to a closed mold. When that scorch time is delayed, the gas evolution may coincide with the crosslink network formation. Furthermore, in addition to these various unexpected processing and engineering advantages, these novel peroxide formulations provide a completely tack free surface when cured in the presence of air.



**TABLE 3**

Run #	1	2	3	4	5	6
Uniroyal X3378 EPDM	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.						
N774 Carbon Black	100	100	100	100	100	100
Sunpar 2280 Oil	30	30	30	30	30	30
ZnO	4	4	4	4	4	4
Stearic acid	0.5	0.5	0.5	0.5	0.5	0.5
AgeRite D	1	1	1	1	1	1
DC40KEP 40% Dicumyl Peroxide	8.0	5.2	5.2	5.2	5.2	5.2
HVA-2	-	2.0	2.0	2.0	2.0	2.0
Sulfads (98%)	-	0.5	0.3	0.3	-	-
VULTAC 5 (75%)	-	-	-	0.5	0.8	0.5
Durax (98%)	-	0.3	0.5	-	-	0.3
MDR at 170°C						
M <sub>H</sub> in-lb	24	30	30	31	32	31
T <sub>S1</sub> min	.53	.83	.76	.69	.55	.54
T <sub>C90</sub> min	7.9	5.9	5.6	5.5	4.9	4.7
Compression set 70 hrs. at 150°C						
% Compression Set	21	30	24	27	22	19
Air cure at 400°F for 10 min.						
Surface	10	1	1	1	1	1
Gassing	No	Yes	Yes	Yes	No	No

Example 4 - Novel peroxide-additive compositions are studied using a fast cure, higher ethylene EPDM (Royalene 509 EPDM) using dicumyl peroxide and HVA-2 in Table 4.

5 The prior art formulations (Table 4, runs #2 and #3) provided a good surface after hot air cure, but unfortunately, resulted in very poor 150°C/70 hour compression set physical properties and exhibited a highly undesirable odor. These 61% and 71% compression set values are similar to what one would obtain from a sulfur vulcanization, not a peroxide cure. In addition, the prior art samples #2 and #3 which used 2.5 parts of Sulfads®, had large gas bubbles trapped at the center of  
10 the cured parts; considered an unacceptable cure defect.

In contrast, all the samples (runs #5 to #12) using the novel peroxide-additive composition taught by this invention, provided outstanding physical properties, i.e., very low compression set values at 150°/70 hours, very similar to that obtained by the use of the peroxide control reaction with no coagent or sulfur  
15 additive (Table 1, run #1). In that control reaction a 15% compression set was obtained, despite the very sticky surface (10 out of 10) for the hot air cure study. The blend of HVA-2 coagent with peroxide (run #4) and the antioxidant AgeRite D (typically incorporated as standard practice in rubber formulations) provided a good compression set value at the reduced 120°C/70 hour test temperature, but  
20 unfortunately, a poor surface (7 out of 10) when crosslinking in the presence of air. All samples included in this invention, runs #5 to #12, provided excellent physical properties associated with a peroxide cure, plus a tack free surface when cured in the presence of air.

TABLE 4

	1	2	3	4	5	6	7	8	9	10	11	12
Royalene 521 EPDM	100	100	100	100	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.												
N550	50	50	50	50	50	50	50	50	50	50	50	50
N 990	25	25	25	25	25	25	25	25	25	25	25	25
LW150 oil	20	20	20	20	20	20	20	20	20	20	20	20
ZnO	4	4	4	4	4	4	4	4	4	4	4	4
Stearic Acid	1	1	1	1	1	1	1	1	1	1	1	1
AgeRite D	1	1	1	1	1	1	1	1	1	1	1	1
DC40 KEP 40%	7.2	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
HVA-2	-	2.2	-	2.2	2.2	2.2	2.2	2.2	2.2	2.2	2.2	2.2
SR-206	-	-	2.2	-	-	-	-	-	-	-	-	-
Sulfads 98%	-	2.5	2.5	-	0.5	0.3	0.3	-	0.3	0.3	0.3	0.3
Vanax A 98%	-	-	-	-	-	-	-	0.3	-	-	-	-
M. Tuads 98%	-	-	-	-	-	0.2	-	-	-	-	-	-
Durax 98%	-	-	-	-	-	-	0.2	0.2	-	-	-	-
Unads 98%	-	-	-	-	-	-	-	-	0.2	-	-	-
M. Zimate 98%	-	-	-	-	-	-	-	-	-	0.2	-	-
M. Niclate 98%	-	-	-	-	-	-	-	-	-	-	0.2	-
M. Cumate 98%	-	-	-	-	-	-	-	-	-	-	-	0.2
MDR at 170°C, 1 degree arc												
M <sub>H</sub>	39	43	27	45	40	38	40	42	38	38	41	37
T <sub>S1</sub>	.45	.68	.75	.38	.60	.62	.58	.42	.65	.46	.56	.53
T <sub>C90</sub>	7.1	4.3	2.5	4.6	4.8	5.3	4.6	4.9	4.8	5.1	4.8	5.2
Cure in air at 400°F for 12 min.												
Surface	10	1	1	7	1	1	1	1	1	1	1	1
Percent Compression set 120°C for 70 hours												
% Set	11	46	-	8	16	18	14	-	15	-	-	-
Percent Compression set 150°C for 70 hours												
% Set	15	61	71	-	-	-	19	17	-	21	17	21

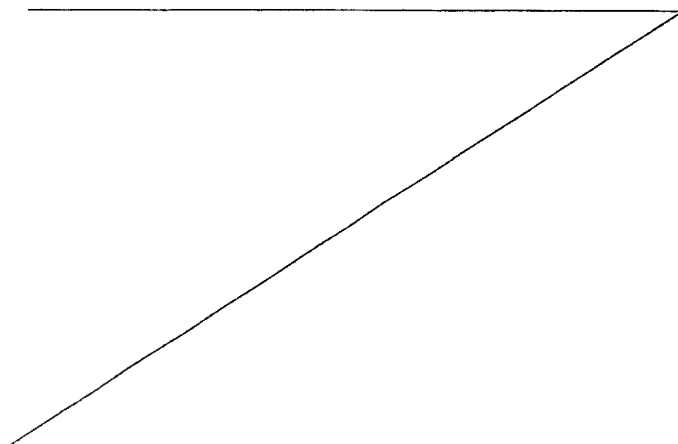
Example 5 - Evaluation of common classes of coagents to demonstrate the

uniqueness of the use of bismaleimide coagents with selected additives, for utility in providing good overall peroxide-type physical properties with the ability to cure the surface of a rubber (tack-free surface) in the presence of air.

5 The bismaleimide is evaluated in combination with Sulfads® and VULTAC® 5 together with dicumyl peroxide for crosslinking a fast cure type EPDM (Royalene® 509 EPDM).

10 The prior art teachings use various coagents both monomeric and non-monomeric to improve crosslinking efficiency, but have no relation to their effect with air contact. We found that the use of these various coagents are just as likely to increase reactions with oxygen and make the rubber surface more tacky after curing in the presence of air, versus the bismaleimide type.

15 In Table 5, the HVA-2 (bismaleimide-type coagent) provided a tack free surface with the blend of 0.3 parts Sulfads and 0.2 parts of VULTAC 5. Use of other agents provided a sticky surface, e.g., TAC (triallyl cyanurate), SR-350 (trimethylolpropane trimethacrylate), 1.2 BR (1,2 liquid butadiene rubber), Santolink XI-100 (allyl glycidyl ether alcohol resin), TAP (triallyl phosphate) and pBQ (para benzoquinone).



**TABLE 5**

Run #	1	2	3	4	5	6	7	8	9
Royalene 509 EPDM	100	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.									
N550	50	50	50	50	50	50	50	50	50
N990	25	25	25	25	25	25	25	25	25
LW150 oil	20	20	20	20	20	20	20	20	20
ZnO	4	4	4	4	4	4	4	4	4
Stearic acid	1	1	1	1	1	1	1	1	1
AgeRite D	1	1	1	1	1	1	1	1	1
DC 40KEP	5.4	5.4	5.4	5.4	5.4	5.4	5.4	5.4	5.4
Sulfads 98%	-	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
VULTAC 5 98%	-	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2
HVA-2	-	-	2.0	-	-	-	-	-	-
TAC	-	-	-	3.0	-	-	-	-	-
SR-350	-	-	-	-	4.0	-	-	-	-
1,2 BR	-	-	-	-	-	6.0	-	-	-
Santolink XI-100	-	-	-	-	-	-	4.0	-	-
TAP	-	-	-	-	-	-	-	4.0	-
pBQ	-	-	-	-	-	-	-	-	1.0
MDR Cure at 170°C									
M <sub>H</sub> in-lb	32	27	42	26	32	27	19	23	22
T <sub>C90</sub> min	7.7	6.6	4.5	7.6	6.7	8.2	7.0	7.2	9.5
Cure in air at 200°C									
Surface	10	4	1	7	7	4	4	4	4



Example 6 - Use of low levels of prior art sulfur accelerators blended with HVA-2 and dicumyl peroxide does not provide tack free surface. Unexpectedly, use of low level prior art compounds with HVA-2 and select compounds provide a tack-free surface when crosslinking a fast cure EPDM (Royalene® 509) in the presence of air.

Using 0.2 parts of Methyl Zimate (98% zinc dimethyldithiocarbamate) or 0.2 parts Unads (98% tetramethylthiuram monosulfide) with HVA-2 provided poor surface cure (sticky surface) when curing in the presence of air. Thus, it is not obvious to use very low levels of the prior art compounds in combination with HVA-2 (N,N' phenylene bismaleimide) coagent and peroxide to provide a tack-free surface for crosslinked elastomers in the presence of air.

Unexpectedly, when VULTAC® 5 (75% alkyl phenol disulfide polymer) or Vanax® A (98% 4,4'dithiodimorpholine) is blended with these low levels of prior art compounds, e.g., Methyl Zimate or Unads, one obtains a tack-free surface when crosslinking in the presence of air.

Thus, a unique blend of peroxide, HVA-2 coagent, and low levels of dithiocarbamates or thiurams with VULTAC 5 or Vanax A provides a crosslinked polymer with a cured surface in the presence of air. In addition, no gassing was noted, thus, solid crosslinked parts can be formed with these novel blends.

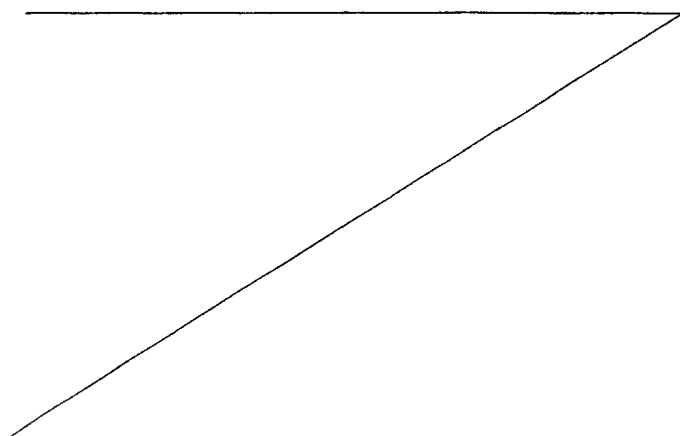


TABLE 6

Run #	1	2	3	4	5	6	7	8	9
Royalene 509 EPDM	100	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.									
N774	75	75	75	75	75	75	75	75	75
Sunpar 2280 Oil	20	20	20	20	20	20	20	20	20
ZnO	5	5	5	5	5	5	5	5	5
Stearic acid	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
DC40KEP 40%	7.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
HVA-2	-	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
M. Zimate 98%	-	0.2	0.2	0.2	0.2	-	-	-	-
Unads 98%	-	-	-	-	-	0.2	0.2	0.2	0.2
Vanax A 98%	-	-	0.5	-	-	-	0.5	-	-
Sulfur	-	-	-	0.15	-	-	-	0.15	-
VULTAC 5 75%	-	-	-	-	0.67	-	-	-	0.67
MDR at 170°C									
M <sub>H</sub> in-lb	43	42	41	43	41	43	40	42	41
T <sub>S1</sub> min	0.38	0.34	0.45	.042	0.40	0.47	0.50	0.47	0.49
T <sub>C90</sub> min	7.2	4.6	4.8	4.4	4.4	4.5	5.3	4.7	4.5
Cure in air at 420°F for 8 min.									
Surface	10	7	1	4	1	7	1	4	1
Gassing	No	No	No	No	No	No	No	No	No

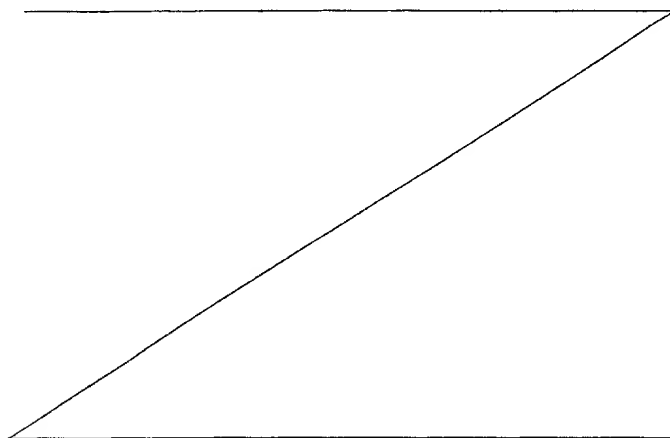
Example 7 - The use of low levels of prior art elemental sulfur with low levels of prior art Sulfads®, with or without optional HVA-2 monomeric coagent, provides good surface cure in hot air, however, poor physical properties (61% and 49% Compression Set).

Thus, the concept of using low levels of known prior art sulfur compounds with suggested coagent and peroxide does not readily lead to a combination of good surface cure and desired physical properties expected.

TABLE 7				
	1	2	3	4
Keltan 4906 EPDM	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.				
N 774	80	80	80	80
N 330	20	20	20	20
LW 150 Oil	40	40	40	40
ZnO	6	6	6	6
Stearic Acid	1	1	1	1
AgeRite D	1	1	1	1
DC40P (40%)	5	5	5	5
HVA-2	-	-	1.8	1.8
Sulfur	-	0.2	-	0.2
Sulfads (98%)	-	0.4	-	0.4
420°F for 8 min. cure in air Surface (1 to 10) 10 = Sticky	10	1	10	1
Compression set 120°C/70 hrs.				
Percent Compression Set	25%	61%	12%	49%

Example 8 - In this example, it is shown that the novel peroxide-additive composition performs well in filled and non-filled EPDM formulations.

Run #2 contains only EPDM and the novel peroxide-additive formulating, i.e., no oil, carbon black, ZnO or stearic acid. Note: the novel formulation, as taught in the practice of the invention, provides a completely tack free surface when cured in the presence of air, (1 out of 10) where 10 is tacky. Addition of normal levels of carbon black and oil (run #4) provides excellent tack free surface in the presence of air. Use of very high levels of carbon black (that exceed the amount of EPDM used) and high levels of oil as shown in Table 8, runs #6 and #8, show the expected reduction in the level of crosslinking ( $M_H$ ) for an organic peroxide type cure. However, the resulting surface cure in the presence of air is surprisingly good (4 out of 10) as shown in run #8, when the level of peroxide-additive blend was increased. It is well known that peroxide cures are adversely affected by high levels of oil and carbon black. The peroxide radicals can be readily abstract hydrogens from the oil as well as from the rubber. When the oil concentration is increased, this significantly reduces the peroxide cure efficiency. This is shown by the reduction in  $M_H$  for the two peroxide controls, with no additives (compare normal levels of oil and carbon black run #3 to high levels of oil and carbon black run #5).



**TABLE 8**

Run #	1	2	3	4	5	6	7	8
Royalene 509 EPDM	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.								
N774 carbon black	-	-	75	75	130	130	130	130
Sunpar 2280 oil	-	-	20	20	60	60	60	60
ZnO	-	-	5.0	5.0	5.0	5.0	5.0	5.0
Stearic acid	-	-	0.5	0.5	0.5	0.5	0.5	0.5
DC40KEP 40%	7.00	5.00	7.00	5.00	7.00	5.00	10.5	7.50
HVA-2	-	1.50	-	1.50	-	1.50	-	2.25
VULTAC 5 75%	-	0.67	-	0.67	-	0.67	-	1.00
B. Zimate 98%	-	0.20	-	0.20	-	0.20	-	0.30
Crosslinking MDR Cure at 170°C, 1° arc								
M <sub>H</sub> in-lb	40	31	43	41	18	24	26	31
T <sub>S1</sub> min	0.45	0.46	0.38	0.40	0.54	0.50	0.43	0.44
T <sub>C90</sub> min	8.3	5.8	7.2	4.4	7.0	4.2	6.6	3.8
Cure in air at 375°F for 14 min.								
Surface	7	1	10	1	10	6	10	4

Example 9 - Novel peroxide-additive blends are used to hot air cure a very highly filled EPDM.

This highly filled EPDM contains a large amount of carbon black and oil. Such formulations are only used for sulfur vulcanization, and are not typically used in peroxide cure. The successful use in these examples shows the unexpected tremendous utility of these novel peroxide-additive blends. In addition, the effectiveness of this system is illustrated by providing examples using four different organic peroxides. These are LUPEROX® 231-XL = 40% 3,3,5-trimethyl-1,1-di(t-butylperoxy)cyclohexane; LUPEROX 230-XL = 40% n-Butyl-4,4-di(t-butylperoxy)valerate; Peroximon DC40KEP = 40% dicumyl peroxide and Peroximon F40KEP 40% di(t-butylperoxy)diisopropylbenzene.

The peroxide controls (Table 9, runs #1, #3, #5 and #7) are all tacky due to an under cured surface when crosslinking in the presence of air. They also contain some gas bubbles in the interior of the sample. The novel peroxide-additive formulations used in this invention provide a fully cured surface (non-tacky) when crosslinking in the presence of air, see Table 9, runs #2, #4, #6 and #8.

Unexpectedly, these formulations cure to a higher level of crosslinking to the point that there was no foaming (gas bubbles). Thus, the novel peroxide formulations provide a desirable, solid, crosslinked part with a non-sticky surface upon hot air cure. The cure times vary because of the different peroxides used, but unexpectedly the surfaces are always tack-free, despite the large amount of filler and oil used.

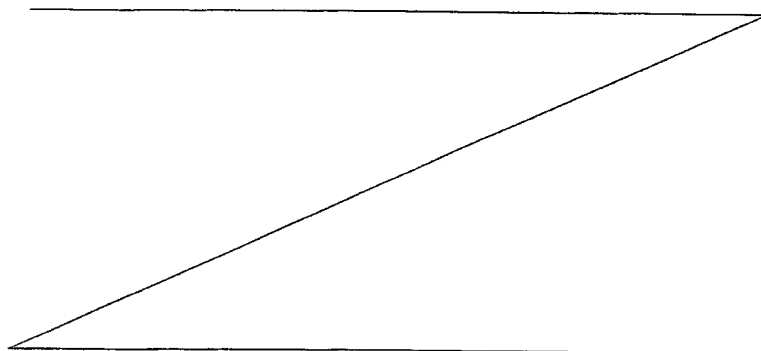


TABLE 9

Run #	1	2	3	4	5	6	7	8
Nordel 2470 EPDM	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.								
N 762	100	100	100	100	100	100	100	100
N 650	100	100	100	100	100	100	100	100
2280 oil	100	100	100	100	100	100	100	100
Lup 231 KE 40%	16.0	10.0	-	-	-	-	-	-
Lup 230 XL 40 %	-	-	16.0	10.0	-	-	-	-
Per DC40P 40%	-	-	-	-	16.0	10.0	-	-
Per F40P 40%	-	-	-	-	-	-	16.0	10.0
HVA-2	-	4.0	-	4.0	-	4.0	-	4.0
Vanax A 98%	-	0.3	-	0.3	-	0.3	-	0.3
Durax 98%	-	0.2	-	0.2	-	0.2	-	0.2
VULTAC 5 75%	-	1.5	-	1.5	-	1.5	-	1.5
MDR Temp.	150°C		165°C		170°C		180°C	
M <sub>H</sub> in-lb	11	26	11	23	14	26	20	29
T <sub>S1</sub> min	.50	.48	.53	.49	.52	.56	.38	.38
T <sub>C90</sub> min	4.6	5.4	4.4	4.5	7.0	5.0	5.4	3.1
Cure in air for 14 min.								
	360°F		390°F		400°F		415°F	
Surface	10	2	10	1	10	1	10	1
Gassing	V.sli	No	Sli.	No	Sli.	No	Sli.	No

Example 10 - ADA (Azodicarbonamide) is a blowing agent which is used to create foamed articles.

In Table 10, ADA is used to produce hot air cured, free-rise, crosslinked EPDM sponge using our novel peroxide-additive formulations to create a tack free surface. The peroxide control with no additive or ADA is provided in Table 10, run #1. Note: the level of crosslinking was  $M_H=19.8$  inch-lbs, as per the Moving Die Rheometer (MDR 2000E, at 200°C and 1° arc). Note, however, the surface was very sticky when crosslinking in the presence of air. The Tangent delta for run #1 (peroxide control) was 0.064, a desirable low value. The MDR can measure  $\tan \delta$  (tangent delta) which can be determined by dividing the viscous moduli by the elastic moduli. A cured compound with a very low tangent delta will be very resilient and exhibit low hysteresis. A higher tangent delta value means that the polymer chains can undergo permanent movement or deformation, after an applied stress. Table 10, run #2, peroxide and ADA with no other additives provided and  $M_H=17.6$  inch-lbs, so the ADA appears to reduce the level of crosslinking, although gas pressure can also affect the reading, compared to run #1 while continuing to provide a sticky surface.

Table 10, run #3, use of peroxide, ADA and coagent blend N,N'-m-phenylene bismaleimide (HVA-2) and 1,3-bis(citraconimidomethyl)benzene (Perkalink 900) provided improved crosslinking performance, based on the higher  $M_H$  of 24.9 in-lbs, but the surface was very sticky (7 rating out of 10) after hot air cure. Coagents with peroxide helps to improve the cure, but continue to provide poor surface due to air inhibition.

Table 10, run #4, low levels of certain additives, ADA and no coagent with peroxide improves the surface (4 to 5 out of 10) after hot air cure, but the crosslinking is very low ( $M_H=14.9$  in-lbs) with an undesirable Tangent Delta of 0.095.

Unexpectedly the use of a coagent blend (HVA-2 and Perkalink 900) with low levels of select additives, peroxide and ADA blowing agent as per our invention provided excellent crosslink density ( $M_H$  of 22.2 in-lbs) and a very good surface (2 out of 10) when cured in hot air (Table 10, run #5). Note: a



uniform foamed EPDM with a density of 30 lbs/cu. ft was produced.

As taught in prior art, Table 10 run #7, use of Vanox ZMT1 (93% Zinc 2-mercapto-toluimidazole) at 5 phr peroxide plus ADA provides a non tacky surface in hot air cure with unfortunately a low  $M_H$  of only 16.9 in-lbs and a

5 corresponding poor tangent delta, compared to run #5.

Use of Captax (98% 2-mercaptobenzothiazole) as per prior art, Table 10 run #8, provides a non-tacky surface as well, however, with further degradation in final crosslinked physical properties ( $M_H$  and tangent delta).

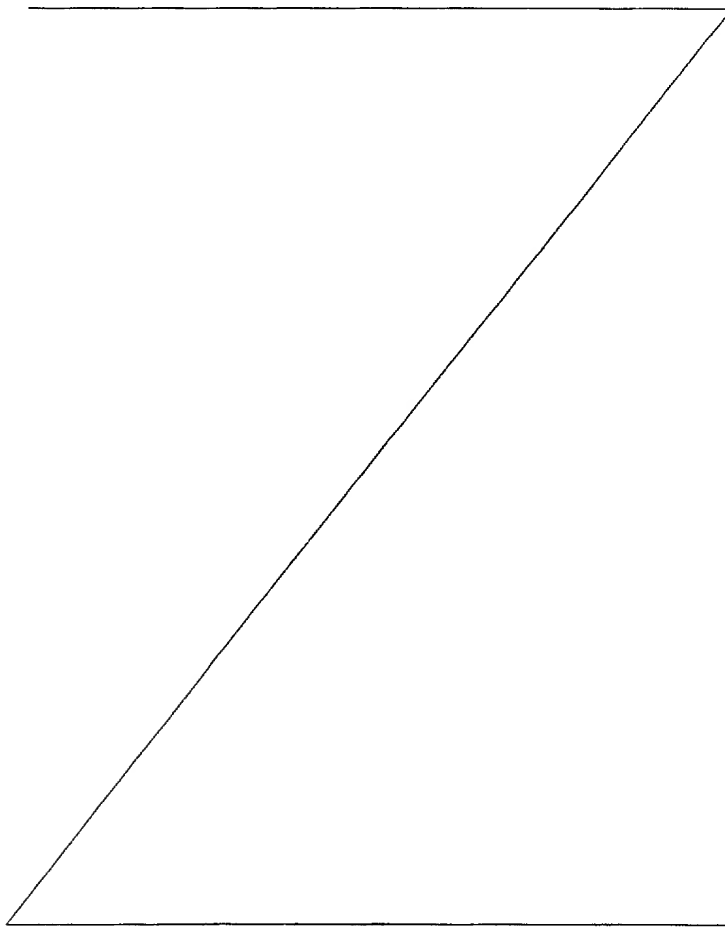


TABLE 10

Run #	1	2	3	4	5	6	7	8
Royalene X3378 EPDM	100	100	100	100	100	100	100	100
The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.								
N774 carbon black	100	100	100	100	100	100	100	100
Sunpar 228 oil	40	40	40	40	40	40	40	40
ZnO	2	2	2	2	2	2	2	2
Stearic acid	2	2	2	2	2	2	2	2
DC40KEP 40%	9.0	9.0	9.0	9.0	9.0	4.5	9.0	9.0
HVA-2	-	-	0.5	-	0.5	2.0	-	-
Perkalink 900	-	-	1.5	-	1.5	-	-	-
Sulfads 98%	-	-	-	0.3	0.3	0.3	-	-
Unads 98%	-	-	-	0.2	0.2	0.2	-	-
Vanox ZMTI 93%	-	-	-	-	-	-	5.0	-
Captax 98%	-	-	-	-	-	-	-	5.0
Azodicarbonamide ADA	-	4.0	4.0	4.0	4.0	4.0	4.0	4.0
Crosslinking in an MDR at 200°C, 1° arc								
M <sub>H</sub> in-lb	19.8	17.6	24.9	14.9	22.2	19.6	16.9	6.7
Tangent Delta	.064	.068	.037	.095	.046	0.64	.093	.247
T <sub>S5</sub> min	0.34	0.36	0.26	0.39	0.33	0.38	0.39	-
T <sub>C90</sub> min	0.78	0.78	0.57	0.69	0.60	0.61	0.75	0.98
Hot Air Cure at 400°F for 7 min.								
Surface	10	5	7	4	1	1	1	4
Density lbs/cu. ft.	74	21	48	22	30	26	29	43
Hot Air cure @ 400°F for 12 min.								
Surface	9	9	9	5	2	1	1	1
Density lbs/cu. ft.	71	24	54	22	30	28	29	41

Example 11 - Common coagents are evaluated by adding them to a sulfur donor - accelerator blend together with LUPEROX® 101-XL [45% 2,5-dimethyl-2,5-di(t-butylperoxy)hexane].

5 These peroxide-coagent-accelerator blends are used to cure standard EPDM to show that the maleimide type coagent, HVA-2 or Vanax MBM is the only one of those tested which is capable of providing the all important combination of good cure with tack-free surface when crosslinking EPDM in the presence of air.

10 Coagents in general are more efficient in Royalene® 521 EPDM compared to Royalene 509 in Example 4/Table 4. Referring to Table 11, run #5 only HVA-2 provides a tack-free surface when used in the practice of our invention, in the presence of air, and has a higher MDR torque than the peroxide control in Table 11, run #1. Although a few of the coagents provide some less reproducible improvement, only the bismaleimide-type coagent, together with the select  
15 additives at low levels, unexpectedly provide an outstanding balance of surface cure and physical properties.

In addition, the previous data in Table 5 and in Table 11 show HVA-2 to have the fastest cure time of all the coagents, while working well with the low levels of additives in the practice of our invention to provide a reduced cure time  
20 in air, a tack free surface with excellent final physical properties. The shortened cure time should help the manufacturer with improved productivity as well as producing a more superior product.

Various Components used in Table 11

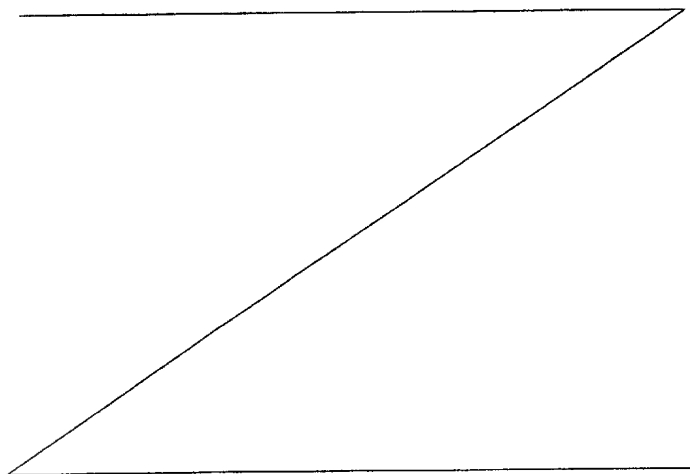
LUPEROX 101-XL (45% 2,5-dimethyl-2,5-di(t-butylperoxy) hexane  
25 Sulfads (98% dipentamethylene thiuram tetra sulfide)  
Santocure TBSI (N-t-butyl-2-benzothiazolesulfenimide)  
TAIC (Triallylisocyanurate)  
SR-350 (trimethylolpropane trimethacrylate)  
HVA-2 (N,N'-m-phenylenebismaleimide)  
30 DAM (diallyl mellitate)

TABLE 11

	1	2	3	4	5	6	7	8	9	10
Royalene 521 EPDM	100	100	100	100	<b>100</b>	100	100	100	100	100
5	The various components below were added as (phr) parts by weight per 100 parts of rubber, "as is," without correcting for assay.									
	N660	80	80	80	80	<b>80</b>	80	80	80	80
	2280 Oil	10	10	10	10	<b>10</b>	10	10	10	10
	Stearic Acid	1	1	1	1	<b>1</b>	1	1	1	1
10	Lup 101XL 45%	4.5	4.5	4.5	4.5	<b>4.5</b>	4.5	4.5	4.5	4.5
	Sulfads 98%	-	0.30	0.30	0.30	<b>0.30</b>	0.30	0.30	0.30	0.30
	TBSI	-	0.15	0.15	0.15	<b>0.15</b>	0.15	0.15	0.15	0.15
	TAIC	-	-	2.00	-	-	-	-	-	-
15	SR-350	-	-	-	2.00	-	-	-	-	-
	HVA-2	-	-	-	-	<b>2.00</b>	-	-	-	-
	ZnDMA	-	-	-	-	-	2.00	-	-	-
	DAM	-	-	-	-	-	-	2.00	-	-
	SR-206	-	-	-	-	-	-	-	2.00	-
20	Perkalink 900	-	-	-	-	-	-	-	-	2.00
	Sulfur	-	-	-	-	-	-	-	-	0.20
	MDR at 185°C									
	M <sub>H</sub>	37	27	32	35	<b>41</b>	27	32	33	31
25	T <sub>S1</sub>	0.34	0.40	0.39	0.41	<b>0.37</b>	0.31	0.40	0.44	0.41
	T <sub>C90</sub>	4.2	3.7	4.3	3.4	<b>2.4</b>	3.7	4.2	4.0	3.6
	Cure in air at 420°F for 8 min.									
	Surface	10	5	4	2	<b>1</b>	1	4	4	4

Example 12 - Various peroxide - sulfur cure blends used for other applications are rated for surface tack (quality) in air cure and percent compression set properties compared to the novel blends of this invention and to standard peroxide and sulfur cures.

5           The novel peroxide cure system in Table 12, run #2 unexpectedly gives an excellent, tack-free surface in air (1 out of 10) with an outstanding compression set equal to the peroxide cures in Table 12, runs #1, #3 and #6. which are very tacky (10 out of 10) after curing in air. Sulfur additives in Table 12, runs #4 and #5, without HVA-2, produces a poor surface with higher, less desirable  
10           compression set valued. A semi EV (semi-efficient sulfur vulcanization system) which uses low levels of elemental sulfur in Table 12, runs #7 and #8, provide expected improved surface (non-tacky) but exhibit the poor compression set values of such EV sulfur cures. The poor compression set values of 77% and 71% for the standard sulfur vulcanization systems, Table 12, runs #9 and #10, provide  
15           good, tack-free surfaces, unlike the peroxide controls of Table 12, runs #1, #3 and #6 which provide excellent 14-16 percent compression set but a very sticky surface.



**TABLE 12**

Comparison of a novel blend (Run #2) of this invention, compared to a various peroxide-sulfur cure systems (Runs #4, #5, #7 and #8), and to standard peroxide (Runs #1, #3 and #6) and standard sulfur cure (Runs #9 and #10).

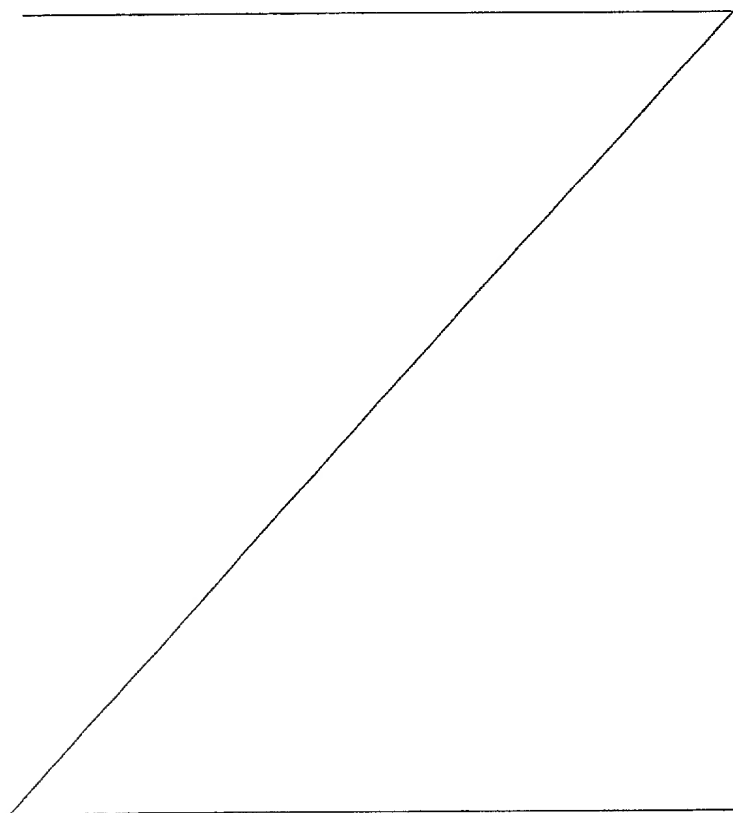
5                      Royalene 521 EPDM                      100  
                          N 660 Black                              75  
                          2280 Oil                                        20  
                          Stearic Acid    1

10	Run #	1	2	3	4	5	6	7	8	9	10
	ZnO	1.00	1.00	1.00	1.00	1.00	3.00	3.00	3.00	3.00	3.00
	DC40KEP	6.00	6.00	7.50	7.50	7.50	8.00	6.00	4.00	-	-
	SR-206	2.00	-	-	-	-	-	-	-	-	-
	HVA-2	-	2.00	-	-	-	-	-	-	-	-
15	Sulfads 98%	-	0.24	-	-	-	-	0.70	1.40	-	2.80
	M. Niclate 98%	-	0.24	-	-	-	-	-	-	-	-
	Altax	-	-	-	-	-	-	0.40	0.80	1.00	1.60
	Unads	-	-	-	-	-	-	0.40	0.80	1.50	1.60
	Durax	-	-	-	0.10	0.20	-	-	-	-	-
20	Sulfur	-	-	-	0.20	0.40	-	0.10	0.20	2.00	0.40
MDR at 170°C											
	M <sub>H</sub> (in-lbs)	26	31	26	25	25	27	21	26	31	29
	T <sub>S1</sub> (mins)	0.74	0.50	0.47	0.45	0.50	0.47	0.85	0.99	1.02	0.93
	T <sub>C90</sub> (mins)	7.1	4.1	7.2	5.3	5.1	7.1	4.3	3.4	3.0	2.6
25	Compression set 70 hrs. at 150°F										
	Percent (%) Compression Set	14	16	16	28	42	15	62	73	77	71
	Cure in air at 400°F for 8 to 12 min.										
	Surface	10	1	10	10	7	10	2	1	1	1

Example 13 - Storage stability of novel peroxide formulations.

The powdered forms of these compositions can be premixed and stored for later use without loss of activity. In Table 13, runs #3 and #5 were Mixture 1 and Mixture 2 respectively, which were aged at room temperature for 3 months.

5 These aged mixtures were compared to freshly prepared peroxide mixtures as shown in Table 13, run #2 and run #4. The rheometer torque versus time curves, and the quality of the crosslinked surfaces which were cured in hot air, are the same for both compounds, within experimental error. MDR torques, times and surface would be closer to the control (run #1) if components reacted during  
10 aging. In the practice of the invention, one can add the various ingredients then add the peroxide, however, premixing the various additives makes it more practical and/or easier to use.



**TABLE 13**

The novel peroxide compositions can be premixed and stored for later use without loss of activity or performance. Two different peroxide formulations were stored for three months.

	<u>Mixture 1</u>	<u>Mixture 2</u>
DC40P	64.6%	64.6%
HVA-2	27.8	22.8
Sulfads	3.8	-
Unads	3.8	-
VULTAC 5	-	8.8
Durax	-	3.8

Keltan 4506 EPDM	100
N 660 Black	50
N 550 Black	50
2280 Oil	35
ZnO	3.0
Stearic Acid	1.5

Run #	1	2	3	4	5
Age of Mixture	0	0	3 mos.	0	3 mos.
DC40P	8.0	-	-	-	-
Mixture 1	-	7.9	-	-	-
Mixture 1	-	-	7.9	-	-
Mixture 2	-	-	-	7.9	-
Mixture 2	-	-	-	-	7.9
Crosslinking Using the MDR @ 170°C, 1° arc					
M <sub>H</sub> (in-lbs)	21.3	25.0	24.6	25.6	25.2
T <sub>S4</sub> (mins)	0.38	0.56	0.55	0.38	0.39
T <sub>C90</sub> (mins)	6.8	4.6	4.5	4.4	4.3
Cure in air at 400°F for 14 min.					
Surface	10	2	1.5	3	2



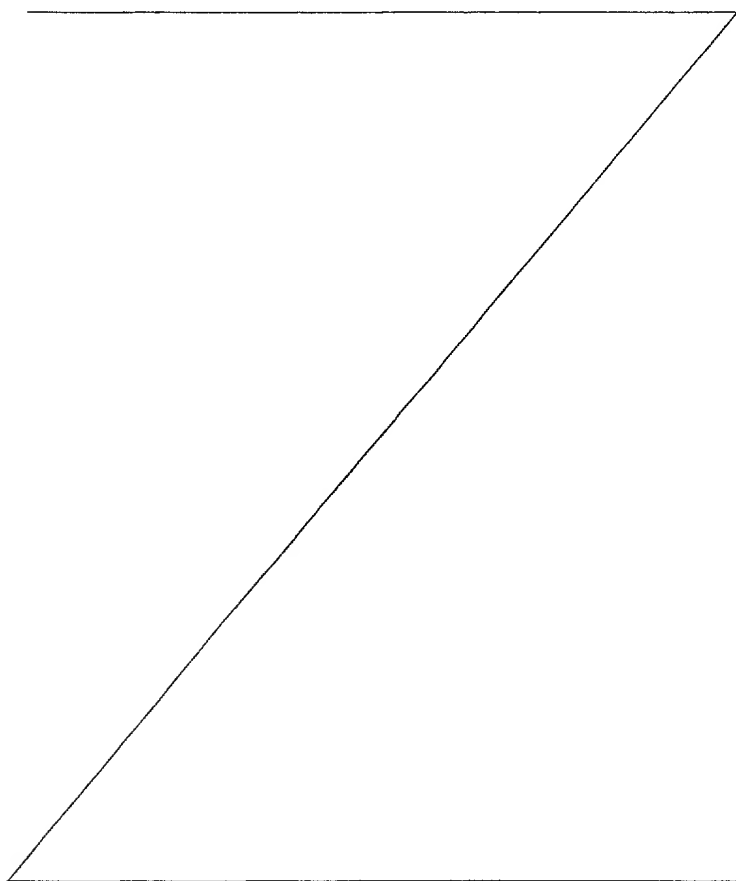
Example 14 - Use of low levels of antioxidants (phenolic, metal dimethyldithiocarbamate type) in combination with bismaleimide coagents and low levels of sulfur donors provide excellent tack-free surfaces when curing elastomers in the presence of air and also provides desirable final physical properties (i.e., low % compression set ).

The novel peroxide air curing formulations which consist of peroxide(s), bismaleimide coagents(s), sulfur donor - sulfur accelerator system can also include low levels of antioxidants. Previously, in Example 4, Table 4, run #11, (M. Niclate, i.e., Methyl Niclate) nickel dimethyldithiocarbamate was used at a low level of 0.2 phr, along with others, e.g., zinc (M. Zimate) and copper (M. Cumate) type dimethyldithiocarbamates along with Sulfads® to successfully produce a tack-free surface with good physical properties (compression set). The Methyl Niclate, whose chemical name is nickel dimethyldithiocarbamate, is sold as antioxidant.

In this present Example 14 and Table 14, a low level of 0.28 phr of Methyl Niclate is used in run #3 compared to a phenolic antioxidant and a sulfenamide accelerator; see run #6 MBPC (2,2'-methylene bis-4-methyl-6-t-butylphenol), and run #7, Santocure TBSI (N-t-butyl-2-benzothiazolesulfenimide). In each case HVA-2 and Sulfads are also included in the formulation, as per the present invention.

Methyl Niclate (nickel dimethyldithiocarbamate) with Sulfads (98% dipentamethylene thiuram tetrasulfide) or Morfax (98% 4-morpholinyl-2-benzothiazole disulfide) is an acceptable combination as per Table 14, runs #2, #3 and #4. Substituting MBPC (2,2'-methylene bis-4-methyl-6-butylphenol) or Santocure TBSI (N-t-butyl-2-benzothiazolesulfenimide) for Niclate (nickel dimethyldithiocarbamate) gives an excellent surface with desirable low compression set physical properties. Thus, low levels of phenolic antioxidants and sulfenamide class of accelerator work well with HVA-2 and low levels of other sulfur accelerators in the novel peroxide compositions of this invention. It is not obvious that low levels of these compounds would be effective, based on the teachings of the prior art. Examples of prior art formulations which use the

acrylic coagent SR-206 (ethylene glycol dimethacrylate) are provided in table 14, run #9 and run #10, where much higher amounts of antioxidants, Vanox MBPC (99% 2,2'-methylene bis-4-methyl-6-t-butylphenol) and Vanox ZMTI (93% Zinc 2-mercapto-toluimidazole) appear to have a poor effect on the surface cure in this  
5 formulation, along with undesirable higher compression set values, particularly for run #10.



**TABLE 14**

Royalene 521 EPDM	100
N 660 Black	75
2280 Oil	20
ZnO	1
Stearic Acid	1

Run #	1	2	3	4	5	6	7	8	9	10
DC40KEP	7.5	6.0	5.0	5.0	5.0	5.0	5.0	6.0	6.0	6.0
SR-206	-	-	-	-	-	-	-	2.0	2.0	2.0
HVA-2	-	2.0	2.0	2.0	2.0	2.0	2.0	-	-	-
Sulfads	-	0.24	0.20	-	-	0.20	0.20	-	-	-
Methyl Niclate	-	0.24	0.28	0.28	-	-	-	-	-	-
Morfax	-	-	-	0.20	0.48	-	-	-	-	-
TBSI	-	-	-	-	-	-	0.28	-	-	-
MBPC	-	-	-	-	-	0.28	-	-	-	2.5
ZMTI	-	-	-	-	-	-	-	-	2.5	-
MDR at 170°C										
M <sub>H</sub>	26	31	29	28	28	29	30	26	26	13
T <sub>S1</sub>	0.47	0.50	0.51	0.42	0.43	0.55	0.54	0.74	0.73	1.44
T <sub>C90</sub>	7.2	4.1	3.7	3.8	3.9	3.9	4.1	7.1	6.7	11.6
Cure in air at 400°F for 12 min.										
Surface	10	1.5	2	2	4	1	1	10	8	10
Compression set 150°C/70 hrs.										
% Comp. Set	16	16	16	15	19	17	19	14	22	49

Example 15 - Crosslinking a fully saturated elastomer (ethylene-octene copolymer).

An ethylene-octene (24%) copolymer G8100 from Dupont Dow, is cured with 2 levels of dicumyl peroxide with and without HVA-2 - VULTAC® 5 - Butyl Zimate in the rheometer and thin samples in hot air. This extended formulation is rubbery, but slightly thermoplastic and not sulfur curable.

In Table 15, run #3 and run #4 are only slightly tacky after curing which is a major improvement because most thermoplastic ethylene polymers would be very sticky in this test, simply because these polymers melt/soften at these temperatures. The oil is added to reduce stiffness but can make the surface more tacky. The oil can be left out at this low filler level. When cooled to room temperature, these samples (runs #3 & 4) were not tacky.

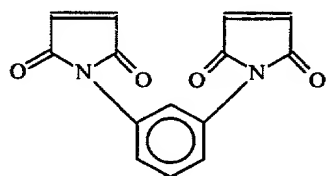
**Table 15**

**Crosslinking a Fully Saturated Elastomer (Ethylene-Octene Copolymer)**

	EG8100	100		
	N 660 black	60		
	2280 oil	5		
	ZnO	3		
	Stearic Acid	1		
	AgeRite MA	0.5		
Run #	1	2	3	4
DC40P	7.0	10.0	4.00	5.72
HVA-2	-	-	1.50	2.14
VULTAC 5	-	-	0.55	0.79
B. Zimate	-	-	0.25	0.36
MDR at 170°C				
M <sub>H</sub> in lb	22	30	30	37
T <sub>S.4</sub> min	0.43	0.39	0.49	0.44
T <sub>C90</sub> min	7.6	6.9	5.3	5.0
Cured in air Surface tackiness rating				
420°F/8 min	10	10	5	5
375°F/18 min	10	10	4	4

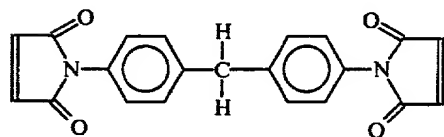
Example 16 - Evaluation of different bismaleimide compounds for use in novel peroxide formulations which can crosslink elastomers with a tack-free surface in the presence of air.

In Table 16, several different bismaleimide compounds were evaluated in the practice of this invention. Unexpectedly, all of these bismaleimide compounds provided good crosslinking efficiency plus tack-free surfaces when tested immediately upon removal from a hot air cure profile of 410°F for eight minutes. The various coagents were evaluated on an equivalent molar bases (taking into account the number of unsaturated functionality) compared to 2 phr of m-phenylene-bismaleimide (MBM). Structures for these various compounds are provided below:



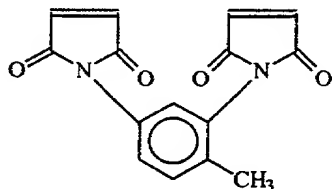
Abbreviation: MBM

N,N'-1,3-phenylene bismaleimide



Abbreviation: MPBM

1-1'-(methylene-di-4,1-phenylene)bismaleimide



Abbreviation: MMBM

N,N'-(4-methyl-1,3-phenylene)bismaleimide

The rubber formulation which was used for these experiments is provided below:

The rubber formulation	Parts
Keltan 2506 EPDM (DSM Copolymer)	100
N660 carbon black (Huber)	75
Sunpar 2280 (Sun Oil)	15
Peroximom DC40KEP	5
Coagent	Molar Equivalent to 2.0 phr of MBM*
Vanax A (®. T. Vanderbilt)	0.2
VULTAC 5 (Elf Atochem)	0.7
Durax (R.T. Vanderbilt)	0.1

\*Taking into account the number of unsaturated groups for each coagent compound.

**TABLE 16**

Run #	1	2	3	4
Coagent Used	None §	MBM	MPBM	MMBM
Crosslinking DSM Copolymer Keltan 2506 EPDM in an MDR 2000E at 170°C, 1° arc				
M <sub>L</sub> (lb-in)	2.270	2.280	2.140	2.180
M <sub>H</sub> (lb-in)	34.90	32.47	27.51	30.08
M <sub>H</sub> -M <sub>L</sub> (min)	32.63	30.19	25.37	27.90
T <sub>S 0.2</sub> (min)	0.310	0.330	0.370	0.345
T <sub>S 0.4</sub> (min)	0.350	0.370	0.430	0.390
T <sub>S 1.0</sub> (min)	0.440	0.465	0.550	0.490
T <sub>C90</sub> (min)	6.850	4.870	5.890	5.260
Crosslinking in a hot air oven at 410°F for 8 minutes				
Surface Tack, 1-10 (10 worst)	10	1	3	2
% Compression Set @ 150°C/70 hours	---	27	46	36

**§Note:** When no coagent was used, 30% more dicumyl peroxide was used in the formulation in an attempt to reduce surface tack. However, even with the increase in peroxide for run #1, surface tack was very poor (10 out of 10).

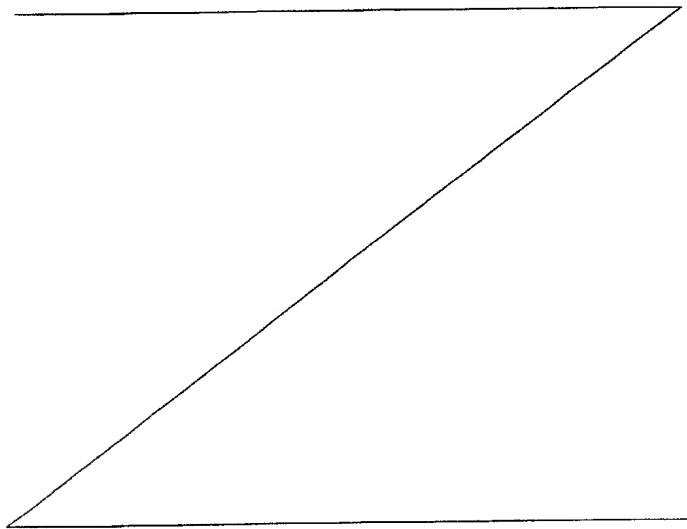
After the MDR2000E testing was completed, the AIR Cure Surface Tack test was performed. All of the samples were heated in the press oven at 410°F for eight minutes. After the eight minutes, the samples were removed and paper towels were pressed onto the hot samples. After cooling, the paper towel was removed at a uniform rate to check for tackiness. Note that after the hot air crosslinked samples had cooled to room temperature, there was very little tack on the surface (except for the control run #1 peroxide cure), with all samples having very similar, low surface tack. All of the samples also displayed their own foaming height, meaning each AIR Cured sample did internally foam to a certain degree. This is partially dependent on the low Mooney Viscosity EPDM used.

Example 17 - Further evaluations of different bismaleimide compounds for use in novel peroxide formulations which can crosslink elastomers with a tack-free surface in the presence of air.

5 The 1,2; 1,3 and 1,4 (ortho, meta, para) isomers of N,N'-phenylene bismaleimide were evaluated for use in novel peroxide compositions which allow crosslinking elastomers in the presence of air and provide a fully cured tack-free surface. The elastomer used in this present example is Nordel®IP EPDM NDR-4640. This is one of the new Dupont Dow metallocene type EPDM elastomers.

10 The various bismaleimide coagents performed well to provide a good tack free surface, as per Table 17, runs #5, #6 and #7, when used with low levels of select additives as per this invention. All of these novel compositions use a unique blend of sulfur containing additives (Sulfads®, VULTAC® 5 and Unads®) at low levels together with the bismaleimide coagent and peroxide.

15 Note that poor surface cure was obtained when the separate components of the peroxide formulations are used individually, see Table 17, runs #1, #2, #3, #4, and #8. Run #1 was the peroxide alone, runs #2 to #4 were the coagents and peroxide, run #8 was the peroxide and sulfur accelerator blend. In each case the surface of these formulations were tacky upon hot air cure at 400°F cure.



**TABLE 17**

Crosslinking Nordel IP NDR-4640 EPDM in Hot Air Oven Evaluation of various bismaleimide compounds in the practice of this invention.

Run #	1	2	3	4	5	6	7	8
DCKEP	7.00	4.20	4.20	4.20	4.20	4.20	4.20	7.00
1,2 PDM	-	2.10	-	-	2.10	-	-	-
1,3 PDM	-	-	2.10	-	-	2.10	-	-
1,4 PDM	-	-	-	2.10	-	-	2.10	-
Sulfads	-	-	-	-	0.28	0.28	0.28	0.28
VULTAC 5	-	-	-	-	0.28	0.28	0.28	0.28
Unads	-	-	-	-	0.14	0.14	0.14	0.14
MDR2000E 175°C, 1°arc								
M <sub>H</sub> (lb-in)	33.8	27.6	37.2	43.3	24.1	34.5	36.2	24.8
T <sub>S,4</sub> (min)	0.29	0.29	0.26	0.27	0.45	0.44	0.44	0.37
T <sub>C90</sub> (min)	4.59	5.26	3.06	2.94	4.59	3.26	5.18	4.22
140°C								
M <sub>L</sub>	3.26	3.28	3.32	3.30	3.28	3.29	3.29	3.15
T <sub>S,4</sub>	2.01	2.04	1.79	1.99	3.37	3.32	3.29	2.67
T <sub>S,1</sub>	3.33	2.91	2.39	2.72	4.85	4.76	4.78	3.97
195°C								
M <sub>H</sub>	30.5	23.5	34.1	39.7	20.5	31.7	34.2	22.7
M <sub>L</sub>	2.47	2.45	2.71	2.63	2.14	2.14	2.16	2.19
T <sub>S,4</sub>	0.19	0.20	0.17	0.17	0.26	0.25	0.25	0.22
T <sub>C90</sub>	1.01	1.08	0.75	0.75	1.00	0.81	1.09	0.96
Hot Air Cure @ 400°F at 12 minutes								
Tacky Surface Rating	10	8	8	8	4	1	3	10
Hardness	60	60	62	65	60	62	62	58

Formulation: Nordel 4640

100

\*Surface Tackiness Ratings

N 774

80

1 no, 4 slight, 7 moderate, 10 yes

N 990

20

Sunpar 2280

25

ZnO

1

Stearic Acid

1

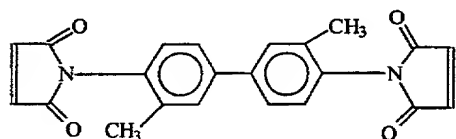
PDM = N,N-(1,2 or 1,3 or 1,4)-phenylenedimaleimide



Example 18 - Further evaluation of various bismaleimide compounds in the practice of this invention.

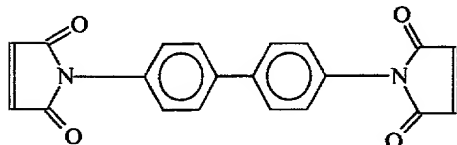
Table 18 shows the results of the evaluation of four coagent bismaleimide compounds for their effectiveness to provide a cured elastomer with tack-free surface upon curing in a hot air oven, when used with peroxides and select low levels of sulfur accelerators, as per the practice of this present invention. In this example, Dupont Dow Nordel IP NDR-4640 metallocene EPDM was chosen as the elastomer for evaluation work.

The monomeric compounds are: 1,1'(3,3-dimethyl-1,1'-biphenyl-4,4'-diyl)-bismaleimide; N,N'-(1,1'-biphenyl-4,4'-bismaleimide; 1,2-bismaleimidoethane and 1,6-bismaleimidohexane. They were compared no N,N'-1,3-phenylenebismaleimide or Vanax MBM, in Table 18. Using our preferred formulation consisting of dicumyl peroxide, Sulfads, VULTAC 5 and Unads, unexpectedly we found that the various bismaleimide coagents provided excellent tack-free surfaces, Table 18, run #2 to run #6, compared to the peroxide control run #1 which provided a very sticky surface.



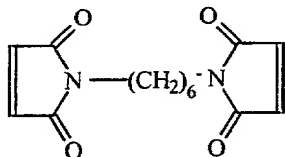
Abbreviation: DMBPDB

1,1'(3,3-dimethyl-1,1'-biphenyl-4,4'-diyl)-bismaleimide



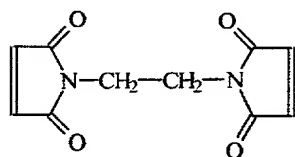
Abbreviation: BPDB

N,N'-(1,1'-biphenyl-4,4'-diyl)-bismaleimide



Abbreviation: BH

1,6-bismaleimidoethane



Abbreviation: BE

1,2-bismaleimidoethane

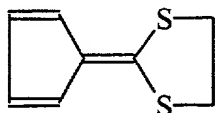
**TABLE 18****Air Cure - Nordel NDR 4640 - Coagent Blend Study**

Run #	1	2	3	4	5	6
DC 40 KEP	7.00	5.00	5.00	5.00	5.00	5.00
Vanax MBM	-	1.50	1.00	1.00	1.00	1.00
DMBPDB	-	-	-	0.25	-	-
BPDP	-	-	-	0.25	-	-
BE	-	-	-	-	0.50	-
BH	-	-	-	-	-	0.50
Sulfads	-	0.28	0.28	0.28	0.28	0.28
VULTAC 5	-	0.48	0.48	0.48	0.48	0.48
Unads	-	0.14	0.14	0.14	0.14	0.14
MDR2000E 175°C						
M <sub>H</sub> (in-lb)	44.0	40.8	36.9	40.1	40.2	39.8
T <sub>S.4</sub> min	0.27	0.38	0.38	0.38	0.39	0.39
T <sub>C90</sub>	4.53	3.05	3.15	3.51	3.20	3.07
MDR 195°C						
M <sub>H</sub>	40.2	38.0	34.3	37.0	36.6	36.9
M <sub>L</sub>	2.94	2.62	2.55	2.60	2.57	2.60
T <sub>S.4</sub>	0.18	0.23	0.23	0.22	0.22	0.23
T <sub>C90</sub>	1.02	0.79	0.80	0.83	0.78	0.79
MDR 140°C						
M <sub>L</sub>	3.85	3.95	3.81	3.91	3.90	3.85
T <sub>S.4</sub>	1.74	2.86	2.85	2.83	2.82	2.79
Evaluation of the surface of crosslinked EPDM, cured in a hot air oven at 400°F for 8 to 12 minutes. Tackiness rating 1 to 10; 10 = sticky surface						
8 minutes	10	2.5	4	2.5	3	4
12 minutes	10	1	2.5	1	1	1
Shore	65	67	66	66	66	66

Formulation: Nordel 4640 100 DMBPDB: 1,1'(3,3-dimethyl-1,1'-biphenyl-4,4'-diyl)-bismaleimide  
N 774 60  
N 990 60 BPDB: N,N'-(1,1'-biphenyl-4,4'-diyl)-bismaleimide  
2280 20 BE: 1,2-bismaleimidoethane  
ZnO 1 BH: 1,6-bismaleimidoethane  
Stearic Acid 1

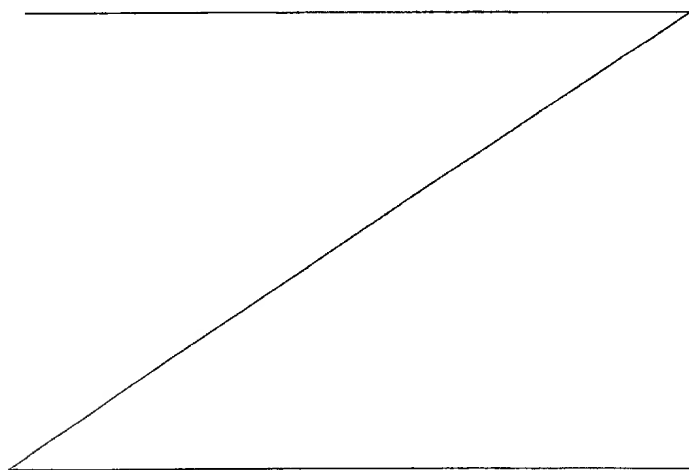
Example 19 - Evaluation of 2-(2,4-cyclopentadien-1-ylidene)-1,3-dithiolane as an additive for use in a novel peroxide formulation, for use in the crosslinking of elastomers in hot air ovens to provide a good surface cure (tack-free surface) with good physical properties.

5 An unsaturated sulfur containing compound was evaluated in combination with a bismaleimide coagent and peroxide. The compound 2-(2,4-cyclopentadien-1-ylidene)-1,3-dithiolane was substituted for the blend of Durax, Vanax A and VULTAC 5 which was used in combination with N,N'-m-phenylene bismaleimide (HVA-2) coagent and dicumyl peroxide. as per the data  
10 in table 19, this unique formulation provided efficient crosslinking with an excellent tack-free surface upon hot air cure at 410°F for eight minutes.



2-(2,4-cyclopentadien-1-ylidene)-1,3-dithiolane

In addition to the good crosslinking performance in the presence of air, thus overcoming the air (oxygen) inhibition, it was found that the final  
15 crosslinked elastomer had excellent final physical properties, based on the desirable, low percent compression set number which was obtained after heat aging the crosslinked EPDM for 70 hours at 150°C.



**TABLE 19**

Run #	1	2
DSM 2506 EPDM	100	100
N660 Carbon black	75	75
Sunpar 2280	15	15
Dicumyl Peroxide 40% on clay	5	5
N,N'-m-phenylene bismaleimide	2	2
Vanax A	0.2	--
VULTAC 5	0.7	--
Durax	0.1	--
2-(2,4-cyclopentadien-1-ylidene)- 1,3-dithiolane	--	1.0
MDR2000E at 170°C, 1° arc		
M <sub>H</sub> - M <sub>L</sub> (in-lbs)	32	30
T <sub>S4</sub> min	0.36	0.34
T <sub>C90</sub>	5.23	5.44
Hot Air Cure at 400°F		
Tacky Surface Rating	2	1
% Compression set, 150°C, 70 hrs.	27%	14%

Example 20 - Use of Polysulfide in combination with select halogenated polymers or silicone rubber, used as additives with peroxide, sulfur accelerators and bismaleimide coagents, to produce a tack-free surface in highly extended EPDM formulations.

5 Nordel IP 4520 (Dupont Dow Elastomers) blended with equal weight of carbon black and high levels of oil would be difficult to cure with ordinary peroxides and is representative of a sulfur cured hose compound formulation. The highly extended EPDM formulations where carbon black and or oil levels are nearly the same as the EPDM usage, represents a formulation which is difficult to cure well  
10 with organic peroxides. When curing a highly filled EPDM in the presence of air, the peroxide radicals will attack the oil and carbon black as if it were rubber, so obtaining a good level of cure and maintaining a tack-free surface in the presence of air is very difficult.

15 Using our novel blend of sulfur accelerators and bismaleimide, we also incorporated small amounts of (FA) polysulfide polymer and (SE6160) silicone rubber as novel additives for use in our invention. We found that the blend of polysulfide polymer and silicone rubber or chlorinated polyethylene or hypalon (chlorosulfonated polyethylene) with peroxide, bismaleimide coagent and low levels of select sulfur accelerators will produce a suitably tack-free cured surface  
20 in the presence of air. So this novel blend provides a means to cure these highly extended EPDM formulation which are commonly cured with sulfur vulcanization. Note that the various sulfur vulcanization systems used in the art, are not sensitive to higher concentrations of oil and carbon black and are capable of fully curing highly extended rubber formulations.

25 Thus we envision that a polymeric masterbatch of peroxide, select sulfur accelerators, bismaleimide coagent, polysulfide could be prepared wherein chlorinated polyethylene and or chlorosulfonated polyethylene could be incorporated as additional additives, as part of our invention.

30 Note the use of silicone rubber without the polysulfide polymer produces a moderately tacky surface. Use of a fluoroelastomer as an additive in our invention in combination with polysulfide does not produce a suitable surface.

## AIR Cure - Nordel IP 4520 - Effect of Polymer Additives

	1	2	3	4	5	6
4520 EPDM	100	100	100	100	100	100
N660	100	100	100	100	100	100
N990	60	60	60	60	60	60
LW150 oil	60	60	60	60	60	60
stearic acid	0.50	0.5	0.50	0.50	0.50	0.50
DCKE 40%	10.0	8.0	8.0	8.0	8.0	8.0
MBM	-	1.35	1.35	1.35	1.35	1.35
Vanax A	-	0.15	0.15	0.15	0.15	0.15
VULTAC 5	-	0.45	0.45	0.45	0.45	0.45
Durax	-	0.05	0.05	0.05	0.05	0.05
Polysulfide	-	-	1	1	1	1
Silicone Rubber	-	4.00	4.00	-	-	-
CPE	-		-	4.00	-	-
Hypalon	-		-	-	4.00	-
Fluoroelas.	-		-	-	-	4.00
RPA 170 C, 0.2° arc, 60 cpm						
Max S' inlb	4.57	5.66	5.82	6.41	7.61	5.94
T <sub>s,5</sub>	0.64	0.46	0.44	0.41	0.35	0.46
T <sub>C90</sub>	7.46	4.36	4.69	4.89	3.88	4.63
Surface	Tacky	MOD	SLI	NO	NO	MOD

Surface:

TACKY - very sticky ; SLI = only slightly tacky; NO = no tack; MOD = moderate tack

Example 21 - In this example we show more data for curing the "difficult to cure by peroxide" highly filled EPDM formulations.

5 In this particular formulation we use more carbon black (a total of 180 parts) than EPDM rubber (used at a total of 100 parts). The use of the novel blend of peroxide, MBM and select sulfur accelerators (listed as blend 1) provides a good surface (run 2) when crosslinking in the presence of air, compared to the standard peroxide (run 1) and at a lower overall curative concentration. Again this is a highly filled EPDM formulation which is very difficult to cure with organic peroxides. One skilled in the art would not consider  
10 such a formulation for use with organic peroxides, and would only use sulfur vulcanization cure systems.

When crosslinking in the presence of hot air, further improvement in surface cure was attained when using a small amount of a chlorosulfonated polyethylene (Hypalon - from Dupont Dow) as a novel additive to be used in the practice of  
15 our invention (run 3). Further improvements in surface cure were noted when the blend of polysulfide and Hypalon (run 4) were used as novel polymeric additives in the practice of our invention.

Use of Chlorosulfonated Polyethylene and Polysulfide as novel additives to crosslink a highly filled EPDM in the presence of air to provide a tack-free surface using organic peroxides

	1	2	3	4
DCKEP	15.00	-	4.00	4.00
Blend 1	-	10.00	8.00	8.00
AgeRite MA	-	-	-	-
MTBHQ	-	-	-	-
* Hypalon 40	-	-	5.00	5.00
Polysulfide	-	-	-	1.00
RPA cure at 170 C, 0.2 degree arc, 60 cpm				
Max S' in-lbs	5.77	6.75	8.72	8.32
T <sub>S5</sub> min	0.42	0.53	0.35	0.35
T <sub>C90</sub> min	6.07	4.41	3.46	3.69
Hot Air Cure at 400 F for 8 to 12 minutes				
Surface Tack:	Tacky	Slight	Very Slight	None

15

## Rubber Formulation

Blend 1

4570 EPDM 80 \*

DCKE 60%

4770 EPDM 20

MBM 27%

N 660 120

Vanax A 3%

N 990 60

VULTAC 5 9%

20

LW 150 80

Durax 1%

Stearic acid 0.8

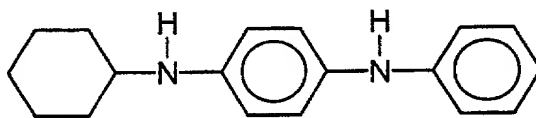
\* a small portion of Hypalon (amount listed above) replaces an equal amount of 4570 EPDM



Example 22- In this example we illustrate several new chemical and polymeric additives which are useful in developing a peroxide formulation for crosslinking elastomers in the presence of hot air.

5 In particular these are: Vanax 6H (N-cyclohexyl-N'-phenyl-p-phenylenediamine) CAS 101-87-1, an antiozonant for diene based polymers; Polysulfide (PTE) and silicone rubber (VMQ).

Vanax 6H (R. T. Vanderbilt)



10 We show that the antiozonant serves as a replacement for the sulfur containing compounds in the practice of this invention. The invention contemplates such substitution of this antioxidant and those of similar structure as full equivalents to the other compounds included within the scope of the definition of compound (B). In addition polysulfide (PTE) also serves this same purpose.

15 Lastly, by an unexpected blend of two polymers PTE and VMQ as polymeric additives one can produce a tack-free surface with peroxides in the presence of hot air. Note no bismaleimide and sulfur compounds or antiozonant additives were used.

TABLE 22

## Crosslinking Royalene 501 EPDM

Use of polysulfide and silicone rubber as well as the use of an antiozonant N-cyclohexyl-N'-phenyl-p-phenylene diamine (Vanax 6H).

## 5 Basic Elastomer Formulation:

100 parts of Royalene 501 EPDM, 70 parts N550 carbon black, 30 parts N-990 carbon black, 20 parts Sunpar LW150 oil, 1 part zinc oxide, 1 part stearic acid

Additive	1	2	3	4	5	6	7
DCP 40KE	7.5	4.6	4.6	4.6	4.6	4.6	7.5
MBM	—	2.2	2.2	2.2	2.2	2.2	—
Vanax A	—	—	0.7	—	—	—	—
Vanax 6H	—	—	—	0.7	—	—	—
Polysulfide	—	—	—	—	0.7	2.2	3.0
VMQ	—	—	—	—	—	—	7.0
Crosslinking using an MDR 2000E 175° C, 1° arc							
M <sub>H</sub> (in-lbs)	35.6	37.8	40.1	36.1	33.3	36.0	27.7
T <sub>S0.4</sub> (min)	0.3	0.26	0.29	0.34	0.26	0.26	0.30
T <sub>C90</sub> (min)	4.39	2.98	3.46	3.63	2.51	2.89	3.98
Hot Air Cure at 400°F (8 to 12 minutes)							
Tacky Surface (tested hot)	Yes	Yes	No	No	No	No	No

Example 23- Examples of using various other dialkyl type peroxides

In this example we crosslink a metallocene based EPDM (Nordel IP 4640) using different organic peroxides. Note that in each case the peroxide without the various additives, was not able to provide a tack-free surface when crosslinking the EPDM elastomer formulation in the presence of hot air. The peroxides listed

5 in Table 23 are:

<u>Peroxide</u>	<u>Chemical Name</u>
Lup 101	2,5-dimethyl-2,5-Di(t-butylperoxy)hexane
MTBPP	4-methyl-4-(t-butylperoxy)-2-pentanone
10 Luperox F/R	Di(t-butylperoxy) diisopropyl benzene
DAPDIB	Di(t-amylperoxy) diisopropyl benzene

This example is provided to illustrate that other dialkyl peroxides can be used in the practice of this invention, besides the most commonly used dialkyl peroxide, dicumyl peroxide, which was extensively evaluated in the previous examples.

## Use of other Dialkyl Peroxides for Hot Air Cure Crosslinking of Nordel IP 4640

EPDM

	1	2	3	4	5	6	7	8
Nordel 4640	100	100	100	100	100	100	100	100
N330	60	60	60	60	60	60	60	60
N990	40	40	40	40	40	40	40	40
LW150	35	35	35	35	35	35	35	35
ZnO	1	1	1	1	1	1	1	1
Stearic Acid	1	1	1	1	1	1	1	1
Lup 101 92.8%	3.60	2.20	-	-	-	-	-	-
MTBPP 84.4%	-	-	4.90	3.00	-	-	-	-
Luperox F/R	-	-	-	-	3.60	2.20	-	-
DAPDIB 79.7%	-	-	-	-	-	-	4.90	3.00
Vanax MBM	-	2.40	-	2.40	-	2.40	-	2.40
Sulfads	-	0.32	-	0.32	-	0.32	-	0.32
Vultac 5	-	0.32	-	0.32	-	0.32	-	0.32
Unads	-	0.16	-	0.16	-	0.16	-	0.16
Crosslinking using an MDR2000E @ 200°C 1° arc								
M <sub>H</sub>	27.4	30.9	21.1	30.7	32.7	34.9	26.3	29.9
T <sub>S4</sub>	0.18	0.24	0.19	0.24	0.17	0.22	0.17	0.21
T <sub>C90</sub>	1.07	0.76	1.16	0.85	0.91	0.72	0.79	0.58
HOT AIR Cure @400°F; Surface Tack Rating								
Tacky Surface ? (Tested hot)	Yes	No	Yes	No	Yes	No	Yes	No

PeroxideChemical Name

Lup 101

2,5-dimethyl-2,5-Di(t-butylperoxy)hexane

MTBPP

4-methyl-4-(t-butylperoxy)-2-pentanone

Luperox F/R

Di(t-butylperoxy) diisopropyl benzene

DAPDIB

Di(t-amylperoxy) diisopropyl benzene

One of skill in the art will also recognize that the compositions of the inventions may also be formulated as masterbatches for convenience in handling and compounding into the polymers for crosslinking in the presence of molecular oxygen. Typical masterbatch carriers include, for example, microcrystalline wax,

polycaprolactone, Ethylene propylene diene monomers polymers (EPDM), ethylene propylene monomers polymers (EPM), ethylene vinyl alcohol polymer (EVA), polyethylene (PE), or mixtures thereof.

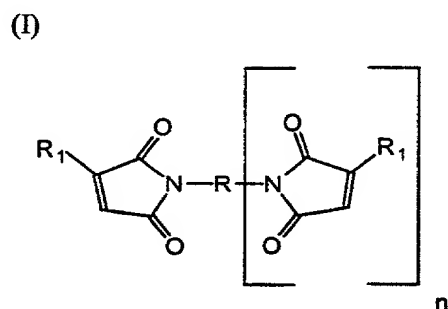
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Algeria	1982	1.00	1000	FAO
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Algeria	1984	1.00	1000	FAO
Algeria	1985	1.00	1000	FAO
Algeria	1986	1.00	1000	FAO
Algeria	1987	1.00	1000	FAO
Algeria	1988	1.00	1000	FAO
Algeria	1989	1.00	1000	FAO
Algeria	1990	1.00	1000	FAO
Algeria	1991	1.00	1000	FAO
Algeria	1992	1.00	1000	FAO
Algeria	1993	1.00	1000	FAO
Algeria	1994	1.00	1000	FAO
Algeria	1995	1.00	1000	FAO
Algeria	1996	1.00	1000	FAO
Algeria	1997	1.00	1000	FAO
Algeria	1998	1.00	1000	FAO
Algeria	1999	1.00	1000	FAO
Algeria	2000	1.00	1000	FAO
Algeria	2001	1.00	1000	FAO
Algeria	2002	1.00	1000	FAO
Algeria	2003	1.00	1000	FAO
Algeria	2004	1.00	1000	FAO
Algeria	2005	1.00	1000	FAO
Algeria	2006	1.00	1000	FAO
Algeria	2007	1.00	1000	FAO
Algeria	2008	1.00	1000	FAO
Algeria	2009	1.00	1000	FAO
Algeria	2010	1.00	1000	FAO
Algeria	2011	1.00	1000	FAO
Algeria	2012	1.00	1000	FAO
Algeria	2013	1.00	1000	FAO
Algeria	2014	1.00	1000	FAO
Algeria	2015	1.00	1000	FAO
Algeria	2016	1.00	1000	FAO
Algeria	2017	1.00	1000	FAO
Algeria	2018	1.00	1000	FAO
Algeria	2019	1.00	1000	FAO
Algeria	2020	1.00	1000	FAO
Algeria	2021	1.00	1000	FAO
Algeria	2022	1.00	1000	FAO
Algeria	2023	1.00	1000	FAO
Algeria	2024	1.00	1000	FAO
Algeria	2025	1.00	1000	FAO
Algeria	2026	1.00	1000	FAO
Algeria	2027	1.00	1000	FAO
Algeria	2028	1.00	1000	FAO
Algeria	2029	1.00	1000	FAO
Algeria	2030	1.00	1000	FAO
Algeria	2031	1.00	1000	FAO
Algeria	2032	1.00	1000	FAO
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Algeria	2036	1.00	1000	FAO
Algeria	2037	1.00	1000	FAO
Algeria	2038	1.00	1000	FAO
Algeria	2039	1.00	1000	FAO
Algeria	2040	1.00	1000	FAO
Algeria	2041	1.00	1000	FAO
Algeria	2042	1.00	1000	FAO
Algeria	2043	1.00	1000	FAO
Algeria	2044	1.00	1000	FAO
Algeria	2045	1.00	1000	FAO
Algeria	2046	1.00	1000	FAO
Algeria	2047	1.00	1000	FAO
Algeria	2048	1.00	1000	FAO
Algeria	2049	1.00	1000	FAO
Algeria	2050	1.00	1000	FAO
Algeria	2051	1.00	1000	FAO
Algeria	2052	1.00	1000	FAO
Algeria	2053	1.00	1000	FAO
Algeria	2054	1.00	1000	

CLAIMS

The subject matter which applicants regard as their invention is particularly pointed out and distinctly claimed as follows:

We Claim:

- 5           1.     A composition comprising:
- a)     At least one compound (A) selected from the group consisting of silicone elastomers and a compound having the formula (I):



Wherein n is 1, or 2 and R is divalent, or trivalent and is selected from the group consisting of acyclic aliphatic groups having from about 2 to 16 carbon atoms, cyclic aliphatic groups having from about 5 to 20 carbon atoms, aromatic groups having from about 6 to 18 carbon atoms and alkyl aromatic groups having from about 7 to 24 carbon atoms, and wherein the divalent, or trivalent groups may contain one or more heteroatoms selected from O, N and S, replacing a carbon atom, or carbon atoms and each R<sub>1</sub> is identical and is hydrogen or an alkyl group of 1 to 18 carbon atoms; and

- b)     At least one compound (B) selected from the group consisting of p-phenylenediamine based antiozonants and sulfur containing organic compounds selected from the group consisting of sulfur containing organic compounds capable of accelerating sulfur vulcanization of a polymer capable of being crosslinked by sulfur, polysulfide polymers and mixtures of said sulfur containing compounds.

2.     A composition as defined in claim 1 additionally containing a free

radical initiator selected from the group consisting of organic peroxides and azo initiators.

3. A composition comprising a composition as defined in Claim 2 and a polymer curable by free radical initiators.
4. A process comprising forming the composition of Claim 3 into a shaped article and then subjecting it in the presence of molecular oxygen to a temperature sufficient to initiate decomposition of the free radical initiator and thereby obtaining a cured shaped article substantially free of surface tack.
5. A composition formed by mixing compounds (A) and (B) as defined in claim one.
6. A composition formed by mixing compounds (A) and (B) as defined in claim 1 and an organic peroxide in any order.
7. A composition as defined in claim 1 wherein compound (A) is selected from bismaleimides and Compound (B) is selected from sulfur accelerators.
8. A composition as defined in claim 1 wherein compound (A) is selected from biscitraconimides and compound (B) is selected from sulfur accelerators.
9. A composition as defined in claim 1 wherein compound (A) is selected from bismaleimides and compound (B) is selected from polysulfide polymers.
10. A composition as defined in claim 1 wherein compound (A) is selected from biscitraconimides and compound (B) is selected from polysulfide polymers.
11. A composition as defined in claim 1 wherein compound (A) is selected from silicone elastomers and compound (B) is selected from polysulfide polymers.
12. A composition as defined in claim 1 also comprising a compound selected from the group consisting of chlorinated polyethylene and chlorosulfonated polyethylene.
13. A composition as defined in claim 2 wherein the free radical initiator is selected from organic peroxides.
14. A surface tack free cured polymer cured in the presence of

molecular oxygen by free radicals generated by decomposition of the free radical initiator in a composition as defined in claim 3.

15. A process for making a surface tack free cured polymer cured by free radicals generated by decomposition of a free radical initiator contained therein in the presence of molecular oxygen which comprises compounding said polymer with a composition as defined in claim 2 and supplying sufficient heat energy to decompose the free radical initiator thus introduced into the polymer.

16. A process for making a curable composition capable of being cured to a tack free surface in the presence of molecular oxygen by a free radical initiator which process comprises compounding a polymer capable of being crosslinked by a free radical initiator with a composition as defined in claim 2.

17. A process for making a curable composition capable of being cured to a tack free surface in the presence of molecular oxygen by a free radical initiator which comprises compounding a polymer capable of being crosslinked by a free radical initiator containing a composition as defined in claim 1 in the presence of a free radical initiator.

18. A composition as defined in claim 1 wherein compound (A) is selected from one or more bismaleimides and compound (B) is selected from the group consisting of dialkylthiuram tetrasulfides, diarylthiuram tetrasulfides, alkylphenol disulfides, tetraalkylthiuram monosulfides, tetraarylthiuram monosulfides and mixtures thereof.

19. A composition as defined in claim 2 wherein the free radical initiator is selected from dialkyl peroxides or peroxyketals.

20. A composition as defined in claim 18 additionally containing a free radical initiator selected from dialkyl peroxides or peroxyketals.

21. A composition as defined in claim 1 wherein compound (A) is selected from one or more bismaleimides and compound (B) is selected from the group consisting of 4,4-dithiomorpholine, acyclicalkyl-2-benzothiazole sulfenamides, cyclicalkyl-2-



benzothiazole sulfenamides, aryl-2-benzothiazole sulfenamides, alkylphenol disulfides and mixtures thereof.

22. A composition as defined in claim 21 additionally containing a free radical initiator selected from dialkyl peroxides, or peroxyketals.
23. A composition as defined in claim 2 comprising dicumylperoxide, N,N-phenylenebismaleimide, 4,4-dithiomorpholine, alkylphenoldisulfide and N-cyclohexyl-2-benzothiazole sulfenamide.
24. A composition as defined in claim 2 comprising dicumylperoxide, N,N-phenylenebismaleimide, dipentamethylene thiuram tetrasulfide, alkylphenoldisulfide and tetramethylthiuram monosulfide.
25. A composition as defined in claim 2 comprising dicumyl peroxide, N,N-phenylenebismaleimide, dipentamethylenethiuram tetrasulfide, alkylphenol disulfide and N-t-butyl-benzothiazole-2-sulfenimide.
26. A composition as defined in claim 1 formulated as a masterbatch on a carrier selected from the group consisting of microcrystalline wax, polycaprolactone, EPDM, EPM, EVA, PE and mixtures thereof.
27. A composition as defined in claim 2 formulated as a masterbatch on a carrier selected from microcrystalline wax, polycaprolactone, EPDM, EPM, EVA, PE and mixtures thereof.

**COMBINED DECLARATION AND POWER OF ATTORNEY**

As a below named inventor, I hereby declare that:

**TYPE OF DECLARATION**

This declaration is of the following type:

- ☒ original
- ☐ design
- ☐ supplemental
- ☐ national state of PCT
- ☐ divisional
- ☐ continuation
- ☐ continuation-in-part (CIP)

**INVENTORSHIP IDENTIFICATION**

My residence, post office address and citizenship are as stated below next to my name, I believe I am the original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

**TITLE OF INVENTION**

**TACK FREE SURFACE CURES OF POLYMERS BY  
ORGANIC PEROXIDES IN THE PRESENCE OF AIR**

## SPECIFICATION IDENTIFICATION

the specification of which:

  X   is attached hereto.

       was filed on                                  as Serial No.  
                                 or Express Mail No., as Serial No. not yet  
known                                  and was amended on  
                                 (if applicable).

       was described and claimed in PCT International Appln.  
No.                                  filed on                                  and as  
amended under PCT Article 19 on                                  (if  
any).

## ACKNOWLEDGEMENT OF REVIEW OF PAPERS AND DUTY OF CANDOR

I hereby state that I have reviewed and understand the  
contents of the above-identified specification, including the  
claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is  
material to the patentability of this application in  
accordance with Title 37, Code of Federal Regulations 1.56.

       In compliance with this duty there is attached an  
information disclosure statement. 37 CFR 1.97.

## PRIORITY CLAIM

I hereby claim foreign priority benefits under Title 35,  
United States Code, 119 of any foreign application(s) for  
patent or inventor's certificate or of any PCT international  
application(s) designating at least one country other than the  
United States of America listed below and have also identified  
below any foreign application(s) for patent or inventor's  
certificate or any PCT international application(s)  
designating at least one country other than the United States  
of America filed by me on the same subject matter having a  
filing date before that of the application(s) of which  
priority is claimed.

  X   no such applications have been filed.

       such applications have been filed as follows.

**COMBINED DECLARATION AND POWER OF ATTORNEY**

As a below named inventor, I hereby declare that:

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This declaration is of the following type:

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of America filed by me on the same subject matter having a  
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priority is claimed.

  X   no such applications have been filed.

       such applications have been filed as follows.

EARLIEST FOREIGN APPLICATION(S), IF ANY FILED WITHIN 12 MONTHS  
(6 MONTHS FOR DESIGN) PRIOR TO THIS U.S. APPLICATION

<u>Country</u>	<u>Application No.</u>	<u>Date of Filing</u>	<u>Priority Claimed Under 37 USC 119</u>
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ALL FOREIGN APPLICATION(S), IF ANY FILED MORE THAN 12 MONTHS  
(6 MONTHS FOR DESIGN) PRIOR TO THIS U.S. APPLICATION

CLAIM FOR BENEFIT OF PRIOR U.S. PROVISIONAL APPLICATION(S)  
(34 U.S.C. §119(e))

I hereby claim the benefit under Title 35, United States  
Codes, §119(e) of any United States provisional application(s)  
listed below:

**PROVISIONAL APPLICATION NUMBER**

**FILING DATE**

60/164,488

November 9, 1999

**POWER OF ATTORNEY**

As a named inventor, I hereby appoint the following  
attorneys to prosecute this application and transact all  
business in the Patent and Trademark Office connected  
therewith.

Stanley A. Marcus, Esq.  
Reg. No. 24,425

William D. Mitchell, Esq.  
Reg. No. 20,977

Royal E. Bright, Esq.  
Reg. No. 26,845

Gilbert W. Rudman, Esq.  
Reg. No. 27,589

Nicholas J. DeBenedictis, Esq.  
Reg. No. 25,670

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Patent Department - 26th Floor  
2000 Market Street  
Philadelphia, PA 19103-3222

Royal E. Bright  
(215) 419-7698

# DECLARATION

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

## SIGNATURE(S)

Full name of sole or first inventor: **Michael F. Novits**

Inventor's signature \_\_\_\_\_

Date \_\_\_\_\_ Country of Citizenship: **U.S.A.**

Residence: **Buffalo, New York, U.S.A.**

Post Office Address: **54 Albemarle Street  
Buffalo, New York 14207  
U.S.A.**

Full name of second joint inventor, if any: **Leonard H. Palys**

Inventor's signature \_\_\_\_\_

Date \_\_\_\_\_ Country of Citizenship: **U.S.A.**

Residence: **Eagle, Pennsylvania, U.S.A.**

Post Office Address: **204 Red Tail Circle  
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U.S.A.**

Full name of third joint inventor, if any: **Gary J. Gullo**

Inventor's signature *Gary J. Gullo*

Date 11-8-00 Country of Citizenship: **U.S.A.**

Residence: **Barrington, New Hampshire, U.S.A.**

Post Office Address: **48 Oak Hill Road  
Barrington, New Hampshire 03825  
U.S.A.**

CHECK PROPER BOX(ES) FOR ANY OF THE FOLLOWING ADDED PAGE(S)  
WHICH FORM A PART OF THIS DECLARATION

- \_\_\_ Signature for fourth and subsequent joint inventors.  
Number of pages added \_\_\_
- \_\_\_ Signature by administrator(trix), executor(trix) or  
legal representative for deceased or  
incapacitated inventor. Number of pages added \_\_.
- \_\_\_ Signature for inventor who refuses to sign or cannot  
be reached by person authorized under 37 CFR 1.47.  
Number of pages added \_\_.
- \_\_\_ Added pages to combined declaration and power of  
attorney for divisional, continuation, or  
continuation-in-part (CIP) application. Number of  
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EARLIEST FOREIGN APPLICATION(S), IF ANY FILED WITHIN 12 MONTHS  
(6 MONTHS FOR DESIGN) PRIOR TO THIS U.S. APPLICATION

<u>Country</u>	<u>Application No.</u>	<u>Date of Filing</u>	Priority Claimed <u>Under 37 USC 119</u>
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As a named inventor, I hereby appoint the following  
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SIGNATURE(S)

Full name of sole or first inventor: Michael F. Novits

Inventor's signature Michael F. Novits

Date 11/07/00 Country of Citizenship: U.S.A.

Residence: Buffalo, New York, U.S.A. NY

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U.S.A.

Full name of second joint inventor, if any: Leonard H. Palys

Inventor's signature Leonard H. Palys

Date 11/07/00 Country of Citizenship: U.S.A.

Residence: Eagle, Pennsylvania, U.S.A. PA

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U.S.A.

Full name of third joint inventor, if any: Gary J. Gullo

Inventor's signature \_\_\_\_\_

Date \_\_\_\_\_ Country of Citizenship: U.S.A.

Residence: Barrington, New Hampshire, U.S.A. NH

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Barrington, New Hampshire 03825  
U.S.A.

CHECK PROPER BOX(ES) FOR ANY OF THE FOLLOWING ADDED PAGE(S)  
WHICH FORM A PART OF THIS DECLARATION

\_\_\_\_ Signature for fourth and subsequent joint inventors.  
Number of pages added \_\_\_\_

\_\_\_\_ Signature by administrator(trix), executor(trix) or  
legal representative for deceased or  
incapacitated inventor. Number of pages added \_\_\_\_.

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